EPA Superfund Record of Decision:

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RECORD OF DECISION

FOR THE

CAROLAWN (OU2) SUPERFUND SITE FORT LAWN, CHESTER COUNTY, SOUTH CAROLINA

SEPTEMBER 1995

PREPARED BY:

U.S. ENVIRONMENTAL PROTECTION AGENCY

REGION IV ATLANTA, GEORGIA

DECLARATION FOR THE RECORD OF DECISIO

SITE NAME AND LOCATION

Carolawn (OU2) Site Fort Lawn, Chester County, South Carolina

STATEMENT OF BASIS AND PURPOSE

This decision document presents the selected remedial action for the Carolawn (OU2) Superfund Site (the Site) located in Fort Lawn, Chester County, South Carolina, which was chosen in accordance with the Comprehensive Environmental Response, Compensation, and Liability Act of 1980 (CERCLA), as amended by the Superfund Amendments and Reauthorization Act of 1986 (SARA), 42 U.S.C. 9601 et seq., and, to the extent practicable, the National Oil and Hazardous Substances Contingency Plan (NCP), 40 C.F.R. Part 300 et seq. This decision is based on the administrative record file for this Site.

The State of South Carolina concurs with the selected remedy.

DESCRIPTION OF THE SELECTED REMEDY

This remedy is the final action for the Site. In the absence of any significant source of contamination in the soil, surface water and sediment at the Site, the No Action alternative was selected as the preferred alternative to address the soil, surface water and sediment. In addition, a groundwater remedy has been selected under

a Record of Decision for Carolawn (OU1). However, should future monitoring of the site (e.g. Five-Year Review) indicate that the site poses an unacceptable risk to the environment, then EPA, in consultation with the State of South Carolina, may initiate clean-up actions under the authority of CERCLA and in accordance with the National Oil and Hazardous Substances Pollution contingency Plan.

STATUTORY DETERMINATIONS

Based on the results of the Remedial Investigation and Risk Assessment conducted for the Carolawn (OU2) Site, EPA has determined that no further action is necessary to ensure the protection of human health and the environment, and the selected remedy is protective of human health and the environment.

Richard D. Green, Associate Director Office of Superfund and Emergency Response Date

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CAROLAWN (OU2) SUPERFUND SITE

FORT LAWN, CHESTER COUNTY, SOUTH CAROLINA

1.0 SITE LOCATION AND DESCRIPTION

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The Carolawn Site, located on approximately 60 acres of land, is an abandoned, waste storage and disposal facility located in Fort Lawn, Chester County, South Carolina. The site is situated less than three miles west of Fort Lawn, and approximately one-half mile south of South Carolina Highway 9 (see Figure 1). Rural and agricultural areas surround much of the site. The Lancaster & Chester Railroad and County Road 841 border the site to the south and Fishing Creek borders the site to the east. Wooded areas and cultivated fields lie to the west and north of the site.

Approximately 30 permanent, single family residences are located north of the site; most of which are situated along South Carolina Highway 9. There are four residences located within 300 yards of the fenced area with a fifth residence located approximately 1,000 yards west of the site.

2.0 SITE HISTORY AND ENFORCEMENT ACTIVITIES

The Carolawn Site was originally owned by the Southeastern Pollution Control Company (SEPCO) of Charlotte, North Carolina. Beginning in 1970, SEPCO used the site as a storage facility for a solvent recovery plant located in Clover, South Carolina. SEPCO

went bankrupt in 1974, and abandoned the Site leaving approximately 2,500 drums of solvents on site. SEPCO had been storing the drummed solvents in anticipation of incinerating the waste. However, neither an incineration permit nor a storage/disposal permit was issued to SEPCO by the South Carolina Department of Health and Environmental Control (SCDHEC).

In January 1975, Columbia Organic Chemical Company (COCC) was contracted to clean up the SEPCO Plant in Clover, South Carolina. As part of this clean up effort, COCC transported and stored the waste of approximately 2,000 drums at the Carolawn Site. As payment for services rendered during the cleanup of the plant in Clover, South Carolina, COCC received the Carolawn property.

After 1975, South Carolina Recycling and Disposal, Inc. (SCRDI), a subsidiary of COCC, controlled the site. During 1978, SCRDI obtained a permit from SCDHEC for a one-time disposal of 300-400 drums containing inert waste. In October 1978 SCRDI was given approval to dispose of empty drums on the 3-acre fenced portion of the property. After the disposal, SCRDI sold the 3-acre fenced area of the site to the Carolawn Company.

In 1978, the Carolawn Company began the construction of two incinerators on the site. With conditional approval of SCDHEC, a

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test burn was conducted with one incinerator; however, full scale incineration never developed. At the time of abandonment of the site by the Carolawn Company, the 3-acre fenced area contained a concrete loading dock, a diked area for storage of tanks and drums, two incinerators, two storage trailers, 14 storage tanks, and as many as 480 drums containing liquid and solid wastes. An additional 660 drums and 11 storage tanks were located outside the fenced area to the north. In 1979, SCRDI was notified by SCDHEC that they would have to clean up the Carolawn site.

During the early 1980's, SCDHEC and EPA conducted site investigations at the Carolawn site. These investigations included collecting environmental and private residential well samples for analysis. The results of these investigations showed the presence of trichloroethane (TCE) and other solvents in nearby residential wells. The results also indicated that the Site was contaminated with high levels of metals and organic compourds. Due to the elevated levels of contamination found and the potential threat for imminent damage to public health and/or the environment, EPA initiated cleanup activities at the Site on December 1, 1981. The cleanup activities continued through February 1982, and included

removal of contaminated soils, drums, and liquid waste from the Site. Subsequently, in December 1982, the Site was proposed for inclusion on the National Priorities List (NPL). The Carolawn Site was finalized on the NPL in September, 1983. Since continued sampling of local residential wells showed persistently high levels of TCE, the Chester Municipal Sewer District's water main from Highway 9 was extended to four of the five residences living near the site. These four residents were connected to this alternative water supply in 1985.

Due to the complexity of the Carolawn Site, and in order to simplify the investigation and response activities, EPA divided the Site into two discrete study areas known as Operable Units (Figure 2). Operable Unit One (OU1) consists of source areas located on a 3-acre parcel within the fenced area of the Site and the groundwater located beneath the entire Site (to include the groundwater beneath Operable Unit Two-OU2). OU2 consists of the land located immediately around the fenced area and the land located north and west of the fenced are (north and west drum areas).

On August 29, 1985, a group of Potentially Responsible Parties (PRPs) (the Carolawn Generators Steering Committee) entered into a Partial Consent Decree with the United States Government to conduct a Remedial Investigation and Feasibility Study (RI/FS) for OU1. The purpose of this RI/FS was to fully characterized the nature and extent of the contamination present at the Site and to identify the relevant alternatives for remedial action. Phase I and Phase II of

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the RI/FS, conducted at the Site between 1985 and 1989, confirmed the presence of volatile organic compounds (VOCS) in the groundwater exceeding Maximum Contaminant Levels ("MCLs") set by the National Primary Drinking Water Regulations in the Safe Drinking Water Act. On September 27, 1989, EPA issued a ROD for OUI which selected a groundwater interception and extraction system as the remedy for groundwater contamination at the Site. It was also determined that due to the effectiveness of the removal actions, no source of contamination remained within the fenced area of the site. However, the findings documented in the ROD for OUI indicated that limited soil data was collected from the west and north drum areas located outside the fence; therefore, collection of additional samples was necessary to confirm the presence or absence of residual soil contamination in these areas.

In response to these concerns, EPA conducted a field investigation

at the Site in 1990. The purpose of the field investigation was to provide additional information on the presence or absence of contaminants in the subsurface soil at the former storage areas situated outside the fenced area. The sampling results indicated the presence of VOCs in the soil. Although this area was addressed during an EPA removal action and again during the 1990 field investigation by the EPA, Environmental Services Division, some uncertainties still existed as to the presence or absence of soil contamination. Based on EPA's review of all the available data, it was determined that a Remedial Investigation and Feasibility Study (RI/FS) needed to be conducted on OU2 in order to develop a baseline risk assessment which would be used to evaluate a final remediation disposition for the OU2 area of concern. Therefore, EPA conducted RI Field activities at the Site in May 1994 and in October 1994.

3.0 HIGHLIGHTS OF COMMUNITY PARTICIPATION

The information repositories, which includes the Administrative Record, were established at the Lancaster County Library in 1989 and the Chester County Library in 1995 and are available to the public at both the information repositories maintained at the Lancaster County Library, 313 South White Street, Lancaster, South Carolina, the Chester County Library, 100 Center Street, Chester, South Carolina and at the EPA, Region IV Library, 345 Courtland Street, Atlanta, Georgia, 30365. The notice of availability of these documents was published in "THE ROCKHILL HERALD", "THE CHESTER NEWS" AND "THE LANCASTER NEWS" on July 24, 1995.

A public comment period for the proposed plan was held from July 24, 1995 to August 24, 1995. A public meeting was held on August 10, 1995, where representatives from EPA answered questions about

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the findings of the RI and the Baseline Risk Assessment and presented EPA's Proposed Plan for the Site.

EPA received oral comments during the August 10, 1995 public meeting, and written comments during the 30 day public comment period. Responses to the comments received by EPA are included in the Responsiveness Summary (Appendix B).

This ROD presents EPA's selected remedial action for the Site, chosen in accordance with CERCLA, as amended by SARA, and to the extent practicable, the NCP. The remedial action selection for this Site is based on information contained in the Administrative Record. The public and State participation requirements under Section 117 of CERCLA, 42 U.S.C. 9617, have been met for this Site.

4.0 SCOPE AND ROLE OF THIS ACTION WITHIN SITE STRATEGY

This ROD addresses the final response action for the Carolawn Site, addressing soil, surface water and sediment. Groundwater has been address under a separate ROD. The Baseline Risk Assessment indicates that no principal threat exists. at the Site, excluding groundwater. The selected alternative in conjunction with the previously selected groundwater remedy, will be protective of human health and the environment and is consistent with the NCP (40 CFR 300. 430(e)).

5.0 SUMMARY OF SITE CHARACTERISTICS

5.1 Climatology

The climate of the area is classified as humid-continental, with long hot summers and short mild winters. The nearest meteorological station is located in Chester, South Carolina, approximately 15 miles from the Site. Examination of Examination of meteorological data over a 30-year period indicate that the mean monthly temperatures range from 42.20F in January to 79.00F in July. The mean annual temperature is 61.10F. The mean annual precipitation is 47.11 inches, which is evenly distributed

5.2 Surface Hydrology

throughout the year.

The topography of the Site is somewhat sloped so rainfall runoff, along with any leached contaminants, would tend to both stand and percolate into the ground and run off into adjacent surface water bodies. There are drainage ditches or drainagepipes which would tend to concentrate and divert runoff directly into adjacent surface water bodies such as Fishing Creek and the Catawba River.

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Fishing Creek is a moderately-sized stream with flow rates of less than 1000 cubic feet per second (cfs). The Catawba River is a moderate to large river with an annual flow rate of 4351 cfs.

5.3 Geologic and Hydrogeologic Setting

The Carolawn Site is located in the eastern Charlotte Belt of the Piedmont Physiographic Province of South Carolina. This belt is characterized by granitoid gneisses with strong compositional layering probably derived from sediments. The bedrock in the vicinity of the Site consists of Lower Metadiorite and Metagabbros. This complex is cut by pegmatite, granite and mafic dikes.

The stratigraphic units encountered at the site during the RI/FS

for OU1 were as follows:

- i) Alluvial deposits;
- ii) Residual and Colluvial clays;
- iii) Residuum and Saprolite; and
- iv) Bedrock.

The upper regions of the bedrock have been altered by in-situ weathering. This weathering has produced a partially to highly decomposed mixture of rock and soil which is referred to as saprolite. Saprolite retains the vestigial mineralogy and structure of the original rock.

The bedrock beneath the Site has undergone several episodes of deformation. These events have created joint and fractures. These structural features influence groundwater flow within the crystalline bedrock. The major structural features noted at the Carolawn site were joints and dikes. Joint measurements revealed the presence of three joint sets with primary sets striking N45øW and N5øW and a minor set striking at N35øW.

All joint sets had vertical to subvertical dips. The mafic dike identified strikes at approximately N45 \emptyset W and is moderately well fractured.

The major hydrostratigraphic unit beneath the Site is the granodiorite bedrock. Saturated conditions were not encountered in the Residuum/Saprolite unit. It may be possible that the Residuum/Saprolite unit may usually be saturated but the RI was conducted during an extended drought and only unsaturated conditions were encountered in this unit. The groundwater in the bedrock is associated with the joints and fractures.

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The actual direction of groundwater flow through the bedrock is dependent upon the orientation of the joints and fractures. The preferred direction of groundwater flow is to the northeast and southeast. Hydraulic data collected during the RI indicates that Fishing Creek is the primary receptor of the groundwater flowing underneath the site. This data also indicates that the mafic dike does not influence, to any great degree, the hydrology of the site.

The estimated groundwater flow velocity is 1.96×10^{-4} centimeters/second (cm/sec). This in equivalent to 0.56 feet/day. Based on this velocity, it would take approximately six years for groundwater originating in the fenced area to reach Fishing Creek.

5.4 Nature and Extent of Contamination

The purpose of the Remedial Investigation (RI) was to gather and analyze sufficient data to characterize the Site in order to perform the Baseline Risk Assessment, which determines the Site's impact on human health and the environment. Both the RI and the Baseline Risk Assessment are used to determine whether remedial action is necessary at the Site.

The RI was designed to focus on the remaining areas of potential contamination not addressed during the RI/FS for Carolawn (OU1). The main portion of the RI was conducted in May 1994. Additional field work was conducted in October 1994.

During this period, samples of soil, surface water and sediment were collected to determine the nature and extent of contamination at the Site. Groundwater was not evaluated in the RI or the Baseline Risk Assessment, since a groundwater remedy addressing all contaminated groundwater at the Site has been selected for the Carolawn (OU1). Contamination at OU2 was characterized by multimedia sampling. Soil (41 surface and 9 subsurface) samples were collected from the area surrounding the three-acre fenced area (see Figure 3). In addition, one surface soil and one subsurface soil sample was collected from an offsite location to establish background conditions for the Site. Four surface water and sediment samples were collected from Fishing Creek, which borders the site to the east (see Figure 4). One of the surface water and sediment samples was collected upgradient of the Carolawn site to established background conditions for the Site. All samples collected during the RI were analyzed for volatile and extractable organic compounds, pesticides, Polychlorinated biphenyls (PCB's) and metals. Additional RI activities included the following: an ecological site reconnaissance of the Carolawn site and the surrounding area was conducted in order to identify the various habitats which are potentially affected by contaminant migration from the Site; an ecological screening to identify endangered and

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threatened species within the site area; and all electromagnetic investigation to locate any buried wastes or metal objects at the site.

Surface Soil Sampling - The sampling results for surface soils are presented in Appendix A. Composite surface soil samples were collected from 41 grids on and around the site (see Figure 3). Purgeable organic compounds were detected in samples from nine of

the grids. Trichloroethylene was detected in sample 7-SLA at a concentration of 27J ug/kg. Tetrachloroethylene was detected in four samples. The concentrations ranged from 3J ug/kg in sample 34-SLA to 10J ug/kg in sample 8-SLA. Toluene was detected in eight samples and ranged in concentration from 2J ug/kc in sample 30-SLA to 25J ug/kg in sample 7-SLA. Purgeable organic compounds were not detected in the background sample, 45-SLA.

The pesticides 4.4 -DDT and 4.4'-DDE were detected in sample 3-SLA at concentrations of 13 ug/kg and 28 ug/kg, respectively. 4.4-DDE was detected in the background sample, 45-SLA, at a concentration of 15J ug/kg.

PCB's were detected in nine samples. PCB-1254 was detected in all nine samples and ranged in concentration from 287 ug/kg in sample 15-SLA to 5,400C ug/kg in sample 1-SLA. Sample 1-SLA also contained 440 ug/kg of PCB-1248 and 700C ug/kg of PCB-1260. PCB's were not detected in the background sample.

Extractable organic compounds were detected in five surface soil samples. Sample 1-SLA contained 4-nitroaniline, fluoranthene, pyrene and chrysene at concentrations of 190J ug/kg, 92J ug/kg, 110J ug/kg and 180J ug/kg, respectively. Sample 15-SLA contained 790J ug/kg of bis (2-ethylhexyl)phthalate and sample 41-SLA contained 3,800J ug/kg of 4-nitroaniline. Extractable organic compounds were not detected in the background sample, 45-SLA.

Presumptive evidence of extractable organic compounds was detected in all the surface soil samples except samples 2-SLA, 26-SLA and 32-SLA. Unidentified extractable organic compounds were detected in all the samples except sample 10-SLA, 32-SLA; 8-SLA and 39-SLA. Sample 4-SLA, 6-SLA and 8-SLA contained the presumptive evidence of petroleum product.

A variety of metals were detected in the surface soil samples including: arsenic, barium, chromium, lead, mercury and magnesium. Elevated concentrations of these metals were detected in one or more samples. Arsenic was detected in most of the samples at concentrations less than 5 mg/kg. The exception was sample 41-SLA which contained 23 mg/kg. Sample 37-SLA also contained arsenic at

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a concentration of 5.7~mg/kg. Arsenic was not detected in the background sample, 45-SLA.

Barium was detected in every sample. With the exception of sample 28-SLA which contained 1,200 mg/kg, the concentrations ranged between 24J mg/kg in sample and 400 mg/kg in sample 5-SLA. Barium was detected at a concentration of 100 mg/kg in the background sample.

Chromium was detected in every sample. Elevated concentrations above background were detected in samples 4-SLA, 5-SLA, 6-SLA, 7-SLA and 41-SLA. The concentrations in these samples ranged from 170 mg/kg in sample 5-SLA to 380 mg/kg in sample 4-SLA. Chromium was detected at a concentration of 14 mg/kg in the background sample.

Lead was detected in all the surface soil samples. Seventeen samples contained concentrations greater than 20 mg/kg. Five samples including: 4-SLA, 5-SLA, 6-SLA, 7-SLA, 14-SLA and 41-SLA contained concentrations greater than 100 mg/kg. Lead was detected at a concentration of 22 mg/kg in the background sample.

Mercury was detected in nine samples including 2-SLA, 4-SLA, 5-SLA, 6-SLA, 7-SLA, 8-SLA, 12-SLA, 14-SLA and 15-SLA. The concentrations ranged from 0.32 mg/kg in the background sample and sample 15-SLA to 1.7 mg/kg in sample 6-SLA.

Magnesium was detected in all the samples. Sample 41-SLA contained an elevated concentration at 26,000 mg/kg. The background sample contained 4,800 mg/kg of magnesium.

Subsurface Soil SamplingÄ Nine subsurface soil samples were collected from grids 1, 3, 5, 11, 13, 15, 26, 33 and 35 (see Figure 3). The analytical results are included in Appendix A. No purgeable organic compounds or pesticides were detected in any of the samples. Sample 5-SLB contained, 48 ug/kg of PCB-1254.

Presumptive evidence of extractable organic compounds was detected in samples 1-SLB, 5-SLB, 15-SLB, 26-SLB, 33-SLB and 35-SLB. The concentrations ranged from 80JN ug/kg of aminoanthracenedione in sample 1-SLA to 4,000JN ug/kg of phenanthrenol in sample 26-SLB. Phenathrenol was also detected at 4,000JN ug/kg in sample 33-SLB. Unidentified compounds were detected in samples 5-SLB, 15-SLB, 26-SLB, 33-SLB and 35-SLB. Sample 5-SLB contained the presumptive evidence of petroleum product. The background sample, 45-SLB, did not contain any extractable organic compounds.

A variety of metals was detected in the subsurface soil samples. Elevated concentrations of magnesium were detected in six of the

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nine samples and ranged in concentration from 6,700~mg/kg in sample 11-SLB to 15,000~mg/kg in sample 3-SLB. The background sample, 45-SLB, contained-3,000~mg/kg of magnesium. An elevated concentration of nickel, 56~mg/kg, was detected in sample 33-SLE. The background sample contained 20~mg/kg of nickel.

Sediment Sampling- Four sediment samples were collected from

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Fishing Creek at the locations indicated on Figure 4. Analytical results are summarized in Appendix A. No purgeable organic compounds, PCB's or pesticides were detected in any of the samples. Two samples contained extractable organic compounds. Sample 2-SD contained one unidentified compound at a concentration of 900J ug/kg. Sample 4-SD contained 15 unidentified compounds and the presumptive evidence of four additional compounds.

A variety of metals was detected in all of the samples. Concentrations of the individual metals were consistent up and down gradient of the site with the exception of sample 3-SD. Arsenic and barium were detected in sample 3-SD at concentrations of 0.91J mg/kg and 24 mg/kg, respectively. Neither of these metals was detected in any other sample.

Surface Water Sampling- Four surface water samples were collected from four locations in Fishing Creek as indicated on Figure 4. Analytical results are summarized in Appendix A. Sample 201-SW is a duplicate of sample 1-SW. No purgeable or extractable organic compounds, pesticides or PCB's were detected in any of the samples. Metals were detected in all of the samples. Primury MCL's were not exceeded for any of the samples. The secondary MCL's for aluminum (0.05-0.2 mg/l), manganese (0.05 mg/l) and iron (0.3 mg/l) were exceeded in all of the samples. The field parameters of pH, specific conductance and temperature were measured at each location. Results are presented in Appendix A.

Ecological Screening - An endangered and threatened species and critical habitat screening was conducted to identify listed species that are found in the Carolawn Site vicinity. Data regarding the actual, past, or potential presence of rare, threatened, and endangered species have been obtained from the United States Fish and Wildlife Service, as well as the South Carolina Department of Natural Resources. Several federally-designated endangered or threatened species are thought to occur in the central and eastern portions of South Carolina. However, there are no critical habitats for federally/state-designated endangered or threatened species on or near the Carolawn site.

Electromagnetic Investigation - The primary purpose of this Electromagnetic investigation (EM) was to locate any buried waste or metal objects at the site. The EM investigatio: was conducted at

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the site using a Geonics EM-31 which is a noncontacting ground conductivity meter. A cartesian coordinate 25 feet by 25 feet grid system was established. Measurements were obtained from the center of each grid. The results of the EM conductivity survey performed at 130 stations are presented as a computer generated contour map on Figure 5.

The data generated consisted mostly of low values ranging from -2 to 98 mmhos/m. The highest value (98 mmhos/m) was due to interference from the fence. Consequently, this value was not used in preparing Figure 5. No magnetic anomalies were detected which would indicate the presence of buried metal objects.

6.0 SUMMARY OF SITE RISKS

A Baseline Risk Assessment was conducted as part of the RI to estimate the health or environmental threats that could result if no further action were taken at the Carolawn (OU2) site. Results are contained in the Final Baseline Risk Assessment Report. A Baseline Risk Assessment represents an evaluation of the risk posed if no remedial action is taken. The assessment considers environmental media and exposure pathways that could result in unacceptable levels of exposure now or in the foreseeable future.

Data collected and analyzed during the RI provided the basis for the risk evaluation. The risk assessment process can be divided into four components: contaminant identification, exposure assessment, toxicity assessment, and risk characterization.

A. HUMAN HEALTH RISK ASSESSMENT

A.1 Contaminant of Concern

Data collected during the RI Were evaluated in the Baseline Risk Assessment. Contaminants were not included in the Baseline Risk Assessment evaluation if any of the following criteria applied:

Inorganic chemicals were eliminated if the maximum detected concentration was less than two times the average background concentration. Organic chemicals were retained regardless of the background concentration because they are not considered to occur naturally.

In absence of Region IV soil screening values, inorganic and organic chemicals were eliminated from further consideration if their maximumdetected concentration did not exceed the EPA Region III screening criteria for residential soil.

EPA Region IV has not developed screening values for sediment ingestion and dermal contact by humans. Therefore, inorganic

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and organic chemicals were eliminated from further consideration if their maximum, detected concentration did not

exceed EPA Region III screening criteria for residential soil.

Chemicals that were retained and evaluated in the Baseline Risk Assessment are known as chemicals of potential concern (COPCs). The following is a summary of the COPCs identified in each media sampled. In addition, a summary table is presented as Table 1 showing all of the COPCs by medium.

Soil. The results of the surfical soil analyses indicated that there are several COPCs present in the soil cover. These compounds include: arsenic, barium, beryllium, calcium, chromium, copper, iron, lead, magnesium, manganese, sodium and polychlorinated biphenyls (PCBs). The results of subsurface soil analyses indicate that there are several COPCs. These compounds include: arsenic, barium, copper, lead, manganese, mercury, tetrachloroethene and toluene. Other concentrations of inorganics and organics were detected in the soil. However, the concentrations of these contaminants were below the typical background concentration ranges for native soils or were below the threshold standards established by EPA.

Surface Water and Sediment. There were no COPCs identified for surface water. In addition, no volatile and extractable organic compounds, pesticides or PCBs were detected in any of the samples. Metals were detected in all of the surface water samples. However, the concentrations of these contaminants were below the typical background concentration ranges.

The sediment analyses revealed that arsenic is the only chemical of potential concern in sediment. In addition, no volatile organic compounds, pesticides or PCBs were detected in any of the samples.

In summary, the results of the Baseline Risk Assessment concluded that there were no chemicals that significantly contributed to the exposure pathways having a Hazard Quotient above 1 or a cancer risk outside of the EPA acceptable range (1E-6 to 1E-4).

A.2 Exposure Assessment

An exposure assessment was conducted to estimate the magnitude of exposure to the contaminants of potential concern at the Site and the pathways through which these exposures could occur. The results of this exposure assessment are combined with chemical-specific toxicity information to characterize potential risks. Human receptors on or near the site were characterized under current and potential future land use (residential) scenarios. The exposure pathways evaluated quantitatively for the current use scenario (for

TABLE 1
HUMAN HEALTH

SUMMARY OF CHEMICALS OF POTENTIAL CONCERN

Chemical Inorganics	Surface Soil	Subsurface Soil	Sediment
Aluminum		X	
Arsenic	X	X	X
Barium	X		
Beryllium	X	X	
Calcium	X	X	
Chromium	X	X	
Copper	X		
Iron		X	
Lead	X	X	
Magnesium	X	X	
Manganese	X	X	
Potassium		X	
Sodium	X		
Vanadium		X	
Pesticides/PCBs			
PCBs	X		

There were no contaminants of potential concern identified for surface

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adults and children) are incidental ingestion of surfical soil, dermal contact with surfical soil, incidental ingestion of sediment from Fishing Creek and dermal contact with sediment in Fishing Creek. The exposure pathways evaluated under the future use scenario, include the four mentioned above as well as incidental ingestion of subsurface soil, and dermal contact with subsurface soil.

water.

After exposure pathways were developed, the concentrations at the exposure points were calculated. These exposure point concentrations were based on the reasonably maximum exposure (RME) scenario - that is, the highest exposure that is reasonably expected to occur at a Site. The RME is calculated by taking the 95% upper confidence limit on the mean of the natural logarithm (ln) transformed data. The data are transformed because the data are assumed to be lognormal.

Once exposure point concentrations were developed, the chemical intake at each exposure point was calculated. These assumptions, along with the exposure point concentrations are used in equations to develop the Chronic Daily Intake (CDI) for each exposure

for each exposure pathway.

A.3 Toxicity Assessment

The purpose of the toxicity assessment is to assign toxicity values (criteria) to each chemical evaluated in the Baseline Risk Assessment. The toxicity values are used in combination with the estimated doses to which a human could be exposed to evaluate the potential human health risks associated with each contaminant. Human health criteria developed by EPA (cancer slope factors and non-cancer reference doses) were preferentially obtained from the Integrated Risk Information System (IRIS, 1993) or the 1992 Health Effects Assessment Summary Tables (HEAST; EPA, 1992). In some cases the Environmental Criteria Assessment Office (ECAO, 1992) was contacted to obtain criteria for chemicals which were not listed in IRIS or HEAST.

Slope factors (SF) have been developed by EPA for estimating excess lifetime cancer risks associated with exposure to potentially carcinogenic contaminants of concern. SFs, which are expressed as risk per milligram per kilogram day, are multiplied by the estimated intake of a potential carcinogen, in mg/kg-day, to provide an upper-bound estimate of the excess lifetime cancer risk associated with exposure at that intake level. The term "upper bound" reflects the conservative estimate of the risks calculated from the SF. Use of this approach makes underestimation of the actual cancer risk highly unlikely. Slope factors are derived from the results of human epidemiological studies or chronic animal

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bioassay data to which mathematical extrapolation from high to low dose, and from animal to human dose, has been applied, and statistics to account for uncertainty have been applied (e.g. to account for the use of animal data to predict effects on humans).

Reference doses (RfDs) have been developed by EPA for indicating the potential for adverse health effects from exposure to the chemicals of concern exhibiting noncarcinogenic effects. RfDs, which are expressed in units of mg/kg-day, are estimates of daily exposure levels for humans, including sensitive subpopulations, that are likely to be without risk of adverse effect. Estimated intakes of contaminants of concern from environmental media (e.g. the amount of a chemical of concern ingested from contaminated drinking water) can be compared to the RfD. RfDs are derived from human epidemiological studies or from animal bioassay data to which uncertainty factors have been applied (e.g., to account for the use of animal data to predict effects on humans).

A.4 Risk Characterization

In this final step of the risk assessment, the results of the exposure and toxicity assessments are combined to provide numerical estimates of the carcinogenic and non-carcinogexlc risks for the Site.

Cancer Risk is expressed as an incremental probability of an individual developing Cancer over a lifetime as a result of exposure to the potential carcinogen. Excess lifetime cancer risks are determined by multiplying the intake level with the slope factor. These risks are probabilities that are generally expressed in scientific notation (1E-06 or lx10-6). An excess lifetime cancer risk of 1E-06 indicates that, as a plausible upper bound, an individual has a one in one million additional chance of developing cancer, over a 70 year lifetime, as a result of site-related exposure to a carcinogen. The NCP states that sites should be remediated to chemical concentrations that correspond to an upper-bound lifetime cancer risk to an individual not exceeding 1E-06 to 1E-04 excess lifetime risk. Carcinogenic risk levels that exceed this range indicate the need for performing remedial action at the site.

The total incremental lifetime cancer risk for offsite residents under current land use conditions was 1E-06. This represents the sum of a child (age 1 to 6), adolescent (age 7-16), and adult (age 7-30), who is exposed to surface soil and sediment. The risk is primarily due to exposure of arsenic in surface soil and sediment. This risk is at the risk level determined to be protective by EPA.

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The incremental cancer risk for future offsite workers was 6E-06. This was the sum of both exposure pathway risks - incidental ingestion of, and dermal contact with, surface soil. The risk was due to incidental ingestion of, and dermal contact with, arsenic, beryllium, and PCBs This risk is within the risk range deemed protective of human health by the EPA.

The lifetime excess cancer risk for future onsite construction workers was 2E-06. This was the sum of all four exposure pathway risks- incidental ingestion of surface and subsurface soil, and dermal contact with surface and subsurface soil. The risk was due to incidental ingestion of, and dermal contact with, arsenic, beryllium, and PCBs (surface soil only) in both surface and subsurface soil. This risk is within the risk range deemed protective of human health by the EPA.

The total incremental lifetime cancer risk for future onsite residents was 2E-05. This was the sum of all four pathway risks - incidental ingestion of soil, dermal contact with soil, incidental ingestion of sediment, and dermal contact with sediment for both child and adult residents. The risk was due to incidental ingestion of, and dermal contact with, arsenic in sediment, and arsenic, beryllium, and PCBs in surface soil.

To characterize potential noncarcinogenic effects, estimated intake levels are compared with toxicity values. Potential concern for noncarcinogenic effects of a single contaminant in a single medium is expressed as a Hazard Quotient (HQ). A Hazard Quotient is calculated for non-carcinogens to assess whether health problems, other than cancer, might be associated with a Superfund site. It is derived by dividing the chemical exposure level at the site by the chemical level determined to be safe. If the Hazard Quotient is greater than 1 there may be concern for potential health effects. Hazard quotients are calculated for each chemical of potential concern found at the site. To assess the overall potential for non-carcinogenic effects Dosed by more than one chemical, all of the hazard quotients calculated for each chemical are added together. The sum of the hazard quotient is called a hazard index (HI). Like the hazard quotient, if the hazard index is greater than 1.0 then the contaminants pose a possible health risk.

An evaluation of the noncarcinogenic risk calculations presented in the risk assessment indicates that all of the hazard indices under the current and future use scenarios are less than 1.0.

The total HI for current adolescent trespassers was 0.03, primarily due to incidental ingestion of, and dermal contact with arsenic, chromium (VI), and PCBs in surface soil.

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The total HI for current offsite child residents (age 1 to 6) was 0.005, due to incidental ingestion of, and dermal contact with, arsenic in surface soil. The total HI for the current off site adult resident was 0.0007, also due to incidental ingestion and dermal contact with arsenic in sediment.

The total HI for future onsite workers was 0.08, primarily due to incidental ingestion of, and dermal contact with PCBs, arsenic, chromium, and manganese in surface soil. Future onsite construction workers exposed to both surface and subsurface soil had a total HI of 0.7, primarily due to incidental ingestion of, and dermal contact with PCBs, chromium, and arsenic in surface soil; and aluminum, arsenic, chromium, and vanadium in subsurface soil.

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The total HI for future onsite child residents(age 1 to 6) was 0.7, primarily due to incidental ingestion of, and dermal contact with, arsenic, chromium, and PCBs in surface soil. The total HI for future onsite adult residents (age 7 to 30) was 0.1, once again primarily due to incidental ingestion of, and dermal contact with, arsenic, chromium, and PCBs in surface soil.

To conclude, carcinogenic risk estimates for current and future conditions are either below the lower limit 1E-6 or within EPA's acceptable range (1E- 6 to 1E- 4). No non- carcinogenic hazard indices exceeded EPA's acceptable level of 1.0. In summary, EPA has determined that risks to human health from contaminants in the soil and sediment are within EPA's acceptable risk range and that remediation of the soil and sediment would not be required for the protection of human health.

B. ECOLOGICAL RISK ASSESSMENT

B.1 Contaminant Identification

A qualitative risk assessment was conducted to determine if ecological chemicals of potential concern (ECOPCs) posed an unacceptable risk to the ecological receptors on and near the Site. ECOPCs are a subset of all chemicals positively identified at the Site. The screening criteria that are used to select ecological chemicals of potential concern are specific to ecological receptors; therefore, ECOPCs may often include different individual chemicals than the human health assessment. The chemicals at the Site were evaluated as follows:

 Chemicals were not listed if they were not detected in the RI environmental samples provided that the sample quantitation limit (SQL) was not in excess of the appropriate screening values;

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- 2) Inorganic chemicals were eliminated if the detected concentrations did not exceed two times the background concentration (provided that the background concentration did not exceed screening levels);
- 3) All chemicals were eliminated if they were only tentatively identified;
- 4) All chemicals with a low frequency of detection (less than 5 percent for any media being evaluated) were eliminated from consideration;
- 5) Chemicals were eliminated from consideration if the maximum

detected concentration did not exceed the appropriate screening value;

6) All inorganic chemicals in surface soils for which the range of detection did not exceed the chemicals natural background concentrations were eliminated from consideration.

The following is a summary of the ECOPCs identified in each media sampled. In addition, a summary table is presented as Table 2 showing all of the ECOPC by medium.

Soil. The results of the surficial soil analyses indicated that there are several ECOPCs present in the soil cover. These compounds include: arsenic, barium, copper, lead, manganese, mercury, zinc, PCBs, tetrachloroethene, and toluene. Other concentrations of inorganics and organics were detected in the soil. However, the concentrations of these contaminants were below the typical background concentration ranges for native soils or were below the threshold standards established by EPA.

Sediment. With the exception of barium, all chemicals detected in sediment were eliminated as an ECOPC. Barium was unable to be eliminated from sediment during the screening process, because no screening value or background concentration was available for this compound. However, barium is not likely to cause a threat to the aquatic environment because it normally precipitates out of solution as an insoluble salt and therefore is less bioavailable to aquatic organisms. It should be noted that it is unlikely that barium in sediment will pose a significant risk to terrestrial organisms at the site. The rationale behind this statement is that it is unlikely that terrestrial organisms will come in direct contact with the sediment at the site. In addition, barium is not known to bioaccumulate; therefore, this limits the possibility that terrestrial as well as aquatic organisms will come into direct contact with these contaminants through the food chain. For these reasons, exposure of terrestrial and aquatic organisms to barium in

TABLE ENVIRONMENTAL HEALTH SUMMARY OF ECOLOGICAL CHEMICALS OF POTENTIAL CONCERN

Chemical Inorganics	Surface Soil	Sediment
Arsenic	X	
Barium	X	X
Copper	X	
Lead	X	
Manganese	X	
Mercury	X	
Zinc	X	

Pesticides/PCBs
PCBs X
Purgeable Organics
Tetrachloroethene X
Toluene X

There were no contaminants of potential concern identified for surface water.

Barium was unable to be eliminated from sediment during the screening process, because no screening value or background concentration was available for this compound. In addition, barium is not known to bioaccumulate; therefore, this limits the possibility that terrestrial as well as aquatic organisms wil come into direct contact with these contaminants through the food chain. For these reasons, exposure of terrestrial and aquatic organisms to barium in sediment was no further evaluated in this Baseline Risk Assessment.

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sediment was not further evaluated in this Baseline Risk Assessment.

B.2 Ecological Exposure Assessment

Once the contaminants have reached the habitat, one or more of three possible exposure routes may come into play for a specific receptor. These exposure routes are 1) ingestion, 2) respiration, and 3) direct contact. Ingestion of Contaminants occurs when an organism ingests contaminated food or incidentally ingests other contaminated media while feeding or through incidental ingestion of contaminated soil. Respiration of contaminants occurs when an organism absorbs contaminants across a respiratory membrane. Contaminants are also absorbed through direct contact with body parts other than the respiratory organs.

In this particular study, the exposure route via ingestion (of soils) was evaluated for the American robin (Turdus migratorius) and the eastern cottontail rabbit (Sylvilagus floridanus) in order to estimate the magnitude of actual or potential exposure to ECOPC in the surface soil. Intake modeling was necessary to estimate the actual dosage of contaminants that these species may be ingesting from the surface soil. Estimates of dosage were based on daily intake rates and the exposure concentration.

Neither the exposure route via respiration or direct contact (dermal) were estimated for terrestrial receptors. The air pathway was not a concern in this particular study and was eliminated. Also, both the inhalation and dermal exposure routes become very

complex to model (EPA, 1993).

The exposure point concentration (EPC) is the concentration of a contaminant in an environmental medium to which a specific receptor is exposed. It is generally calculated using statistical methodology from a set of data derived from environmental sampling. The specific methodology used to derive the exposure point concentrations in this Baseline Ecological Risk Assessment (BERA) is presented below.

For ECOPC and media in which the number of samples was less than 3, the maximum concentration detected was used to represent the exposure point concentration.

For chemicals and media in which the number of samples was equal to or greater than 3, the upper 95 percent confidence limit (UCL) of the log normal arithmetic mean was used to represent the exposure point concentration. In calculating the UCL, one-half the value of the detection limit was used in calculating the log normal mean for all non-detect samples.

Record of Decisio Carolawn (OU2 Superfund Site

For chemicals and media in which the UCL exceeded the $\max/3$ num detected concentration, the maximumconcentra; ion detected was used to represent the exposure point concentration.

In this particular study, the two surrogate terrestrial receptors (American robin and eastern cottontail rabbit) cho en for study are thought to be exposed to contaminated surface oils via either incidental ingestion of the soil or by ingestior of contaminated food. Total exposure of these organisms to the conlaminated surface soil was estimated by approximating how much of the contaminated media and/or food the receptor is taking in on a d Lily basis. This value is otherwise k~own as the daily intake (DI) dose. The equation and process used to calculate the DI lose for each ,of these species is presented in the Baseline Risk ssessment.

B.3 Ecological Toxicity Assessment

The ecological toxicity assessment involves detelmining the types of adverse effects associated with contaminant exposures, the relationship between the magnitude of exposure and adverse effects, and the related uncertainties involved with the assessment. Environmental toxicity data often comes in the form of the concentration or dose necessary in order to induce some observed effect or response. Quite frequently the observed effect is some sort of mortality event such as the death of 5 percent of the population in an experimental environment (i.e. LC or LD50). In the case of this ecological risk assessment, environmental toxicity

data often comes in the form of environmental benchmarks, such as NOAELs or LOAELs, obtained from various research studies.

The Toxicity Values for the ECOPC contained in surface soil that were used to gage relative risk in this BERA were obtained either directly from the literature, from chemical specific documents issued by the Agency of Toxic Substances and Disease Registry, biological reports issued by the United States Fish and Wildlife Service, from chronic No-Observed-Adverse-Effect Level (NOAEL) or chronic Lowest-Observed-Adverse-Effect-Level (LOAEL) obtained from HEAST, March 1994, or Toxicological Benchmarks for Wildlife.

A safety factor of 10 was applied when converting from a chronic LOAEL to a chronic NOAEL. A listing of TRVs for the American robin and the eastern cottontail rabbit for each ECOPC in the surface soil is presented in Tables 3 and 4, respectively.

B.4 Ecological Risk Characterization

Risk Characterization is the final phase of the risk assessment. It is at this phase that the likelihood of adverse effects occurring as a result of contaminant exposure to a contaminant is evaluated In order to give "risk" a numerical value, a Hazard Quotient (HQ) for each ECOPC is-developed.

TABLE

TRVs FOR THE AMERICAN ROBIN

CAROLAWN SITE (OU2) ECOLOGICAL RISK

ASSESSMENT

FORT LAWN, SOUTH CAROLINA

	TRV
CHEMICAL	DERIVATION

LOAEL VALUE	NOAEL VALUE	ROBIN	ROBIN
		SPECIES/REFERENCE	
mg/kg/day	mg/kg/day	LOAEL TRV	NOAEL TRV
INORGANICS			
ARSENIC		Brown-headed cowbird (2)	
1.10E+02	1.10E+01	1.10E+02	
1.10E+01			
COPPER		1-day old chicks (3)	
3.32E+02	3.32E+01	3.32E+02	
3.32E+01			
LEAD		American Kestrel (2)	
5.00E+02	5.00E+01	5.00E+02	
5.00E+01			
MANGANESE		New Hampshire chicks (4)	
7.21E+02	7.21E+01	7.20E+02	
7.20E+01			
MERCURY		Ring-necked pheasant (2)	
4.20E+01	4.20E+01	4.20E+01	
4.20E+00			

ZINC		Domestic hen (2)	
2.03E+04	2.03E+03	2.03 E+04	
2.03E+03			
ORGANICS			
TETRACHLOROETHENE		Mouse (1)	
1.40E+02	1.40E+01	1.40E+02	1.
TOLUENE		Mouse (3)	
2.60E+02	2.60E+01	2.60E+02	
2.60E+01			
PESTICIDES/PCBs			
PCBs		Ring-necked Pheasant (3)	
1.80E+00	1.80E-01	1.80E+00	1.80E-
01			

- (I) HEAST, March, 1994
- (2) Eisler, January, 1988; April, 1988; April 1987; April, 1993
- (3) Opresko, D.M.; B.E. Sample; G.W. Suter II. Toxicological Benchmarks for Wildlife: 1994 Revision
- (4) Gallup, Willis D. and L.C. Norris

A safety factor of IO was applied to the LOAEL value to extrapolate to a NOAEL value.

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TETRACHLOROETHENE

1.40E+01

TABLE 4 TRVs FOR THE EASTERN COTTONTAIL RABBIT CAROLAWN SITE (OU2) ECOLOGICAL RISK ASSESSMENT FORT LAWN, SOUTH CAROLINA

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			TRV
CHEMICAL			
DERIVATION			
		LOAEL VALUE	NOAEL
VALUE	RABBIT	RABBIT	
	SPECIES/REFERENCE	mg/kg/day	
mg/kg/day	LOAEL TRV	NOAEL TRV	
INORGANI	CCS		
ARSENIC	Mouse (3)	1.26E+00	
1.26E-01	1.26E+00	1.26E-01	
COPPER	Mink (3)	1.17E+02	
1.17E+01	1.17E+02	1.17E+01	
LEAD	Rat (3)	8.80E+01	
8.00E+00	8.00E+01	8.00E+00	
MANGANESE	Rat (3)	8.00E+02	
8.80E+01	8.80E+02	8.80E+01	
MERCURY	Mouse (3)	1.32E+02	
1.32E+01	1.32E+02	1.32E+01	,
ZINC	Rat (3)	1.60E+03	
1.60E+02	1.60E+03	1.60E+02	
ORGANIC	2S		

Mouse (2)

1.40E+02

1.40E+02 ú

1.40E+01

2.6 2.60E+01 TOLUENE Mouse (3) 2.60E+02

2.60E+O2 2.60E+01

PESTICIDES/PCBs

PCBs Rat (1) 1.00E+01

1.00E+00 1.00E+01 1.00E+00

TRV- Toxicity Reference Value

- (1) Agency of Toxic Substances and Disease Registry (ATSDR)
- (2) HEAST, March, 1994
- (3) Opresko, D.M.; B.E. Sample; G.W. Suter II. Toxicological Benchmarks for Wildlife: 1994 Revision

A safety factor of 10 was applied to the LOAEL value to extrapolate to a NOAEL value.

Record of Decisio Carolawn (OU2) Superfund Site

The Hazard Quotient (HQ) method was used to define potential risk to the two representative terrestrial receptors via the soil exposure pathway. This method involves: 1) Estimating the exposure of each receptor species to ECOPCs by ingestion of contaminated food and/or soil; 2) Determining from past scientific studies the highest exposure level which produces no observed adverse effects (NOAEL) and the lowest exposure level which produces observed adverse effects (LOAEL) in the representative species; and, 3) Dividing the estimated receptor species exposure level by the NOAEL and LOAEL. A LOAEL based HQ greater than 1 is indicative that there may be a potential for adverse effects on the receptor species.

Using the american robin as a potential receptor for the soll exposure pathway, the LOAEL HQ values ranged from 4.7E-06 to 6.1E-01 and the NOAEL HQ values ranged from 4.7E-05 to 6.1E+00 (See Table 5). In accordance with EPA's draft guidance (Ecological Risk Assessment Guidance for Superfund - Process for Designing and Conducting Ecological Risk Assessments) for Ecological Risk Assessments, remedial goals for the protection of ecological receptors should be bounded by the NOAEL value on the lower end and the LOAEL value on the upper end. Thus, the risk range is between 6.1E-01 to 4.7E-05 which does not exceed EPA's acceptable level of 1.0.

Using the eastern cottontail rabbit, a potential receptor for the soil exposure pathway, the LOAEL values ranged from 8.8E-08 to 6.2E-03 and the NOAEL values ranged from 8.8E-07 to 6.2E-02 (See Table 6). In accordance with EPA's guidance for Ecological Risk Assessments, remedial goals for the protection of ecological receptors should be bounded by the NOAEL value on the lower end and the LOAEL value on the upper end. Thus, the risk range is between 6.2E-03 to 8.8E-07 which does not exceed EPA's acceptable level of 1.0.

In summary, EPA has determined that risks to the ecological receptors from contaminants in the soil are below EPA's acceptable risk range and that remediation of the soil would not be required for the protection of the environment.

7.0 DESCRIPTION OF "NO ACTION" SELECTED ALTERNATIVES

EPA has determined, based on the results of the Remedial Investigation and the Baseline Risk Assessment, that no action is needed for the soil, surface water or sediment. In addition, a groundwater remedy has been selected under a Record of Decision issued for Carolawn (OU1). However, should future monitoring of the site (e.g. Five-Year Review) indicate that the site poses an unacceptable risk to the environment, then EPA, in consultation with the State of South Carolina, may initiate clean-up actions under the authority of CERCLA and in accordance with the National Oil and Hazardous substances Pollution Contingency Plan.

TABLE
SURFACE SOIL HAZARD QUOTIENTS FOR THE AMERICAN ROBIN
CAROLAWN SITE (OU2) ECOLOGICAL RISK ASSESSMENT
FORT LAWN, SOUTH CAROLINA

		DOSE	DOSE	DOSE	LOAEL	NOAEL
HQ	HQ	HQ	HQ	HQ	HQ	
	CHEMICAL	MEAN	MAXIMUM	UCL	TRV	TRV
MEAN+	MAX+	UCL+	MEAN	MAX*	UCL*	
INORG	ANICS					
ARSENIC		9.06E-01	7.63E+00	1.05E+00	1.10E+02	1.10E+01
8.2E-03	6.9E-02	9.5E-03	8.2E-02	6.9E-01	9.5E-02	
COPPER		7.22E+00	5.19E+01	1.01E+01	3.32E+02	3.32E+01
2.2E-02	1.6E-01	3.0E-02	2.2E-01	1.6E+00	3.0E-01	
LEAD		1.16E+01	7.41E+01	1.81E+01	5.00E+02	5.00E+01
2.3E-02	1.5E-01	3.6E-02	2.3E-01	1.5E+00	3.6E-01	
MANGANESE		4.44E+01	1.64E+02	6.83E+01	7.21E+02	7.21E+01
6.2E-02	2.3E-01	9.5E-02	6.2E-01	2.3E+00	9.5E-01	
MERCURY		7.37E-02	5.64E-01	9.13E-02	4.20E+01	4.20E+00
1.8E-03	1.3E-02	2.2E-03	1.8E-02	1.3E-01	2.2E-02	
ZINC		8.40E+01	3.99E+02	1.01E+02	2.03E+04	2.03E+03
4.1E-03	2.0E-02	5.0E-03	4.1E-02	2.0E-01	5.0E-02	
ORGAN	ICS					
TETRACHLOR	OETHENE	9.13E-04	1.56E-03	9.54E-04	1.40E+02	1.40E+01
6.5E-06	1.1E-05	6.8E-06	6.5E-05	1.1E-04	6.8E-05	
TOLUENE		1.22E-03	4.76E-03	1.32E-03	2.60E+02	2.60E+01
4.7E-06	1.8E-05	5.1E-06	4.7E-05	1.8E-04	5.1E-05	
PESTICID	ES/PCBs					
PCBS		9.87E-03	1.09E+00	6.12E-O2	1.80E+00	1.80E-01
5.5E-03	6.1E-01	3.4E-02	5.5E-02	6.1E+00	3.4E-01	
						_

⁺Hazard quotients derived From I.C)AJ~L TRVs

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^{*}Hazard quotienta derived fi'orn NOAJ~L TRVs

EPC - Exposure Point Concentrations

TRV * Toxicity Reference Values

HQ - Hazard Quotient

TABLE SURFACE SOIL HAZARD QUOTIENTS FOR THE

EASTERN COTTONTAIL RABBIT

CAROLAWN SITE (OU2) ECOLOGICAL

FORT LAWN, SOUTH

RISK ASSESSMENT

CAROLINA

		DOSE	Г	OOSE	DOSE	LOAEL
NOAEL	HQ	HQ	HQ	HQ	H	Q HQ
CHEMICA	L	MEAN	MA	XIMUM	UCL	TRV
TRV	MEAN+	MAX+	UCL+	MEAN*	MAX	* UCL*
INORGANICS						
ARSENIC		9.34E-04	7.8	36E-03	1.08E-03	1.26E+00
1.26E-01	7.4E-04	6.2E-03	8.6E-04	7.4E-03	6.	2E-02 8.6E-03
COPPER		6.50E-02	4.6	57E-01	9.12E-02	1.17E+02
1.17E+01	5.6E-04	4.0E-03	7.8E-04	5.6E-01	3 4.	0E-02 7.8E-03
LEAD		2.39E-02	1.5	3e-01	3.73E-02	8.00E+01
8.00E+00	3.0E-O4	1.9E-03	4.7E-04	3.0E-03	1.	9E-02 4.7E-03
MANGANESE		1.45E-01	5.3	37E-01	2.23E-01	8.80E+02
8.80E+00	1.6E-04	6.1E-04	2.5E-04	1.6E-03	6.	1E-03 2.5E-03
MERCURY		2.17E-04	1.6	6E-03	2.69E-04	1.32E+02
1.32E+01	1.6E-06	1.3E-05	2.0E-06	1.6E-0	5 1.	3E-04 2.0E-05
ZINC		8.92E-02	4.2	23E-01	1.07E-01	1.60E+03
1.60E+02	5.6E-05	2.6E-04	6.7E-05	5.6E-0	2.	6E-03 6.7E-04
ORGANIC						
TETRACHLOEOE'	THENE	2.09E-05	3.5	8E-05	2.19E-05	1.40E+02
1.40E+01	1.5E-07	2.6E-07	1.6E-07	1.5E-0	5 2.	6E-06 1.6E-06
TOLUENE		2.29E-05	8.9	06E-05	2.48E-05	2.60E+02
2.60E+01	8.8E-08	3.4E-07	9.5E-08	8.8E-0	7 3.	4E-06 9.5E-07
PESTICIDES/P	CBs					
PCBs		1.75E-04	1.9	3E-02	1.09E-03	1.00E+01
1.00E+00	1.8E-05	1.9E-03	1.1E-04	1.8E-0	1.	9E-02 1.1E-03

⁺Hazard quotients derived from LOAEL TRVs $\,$

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^{*}Hazard quotients derived from NOAEL TRVs

EPC - Exposure Point Concentration

TRV - Toxicity Reference Values

HQ - Hazard Quotient

8.0 DOCUMENTATION OF SIGNIFICANT CHANGES

The selected remedy as presented in this decision document has no difference, significant or otherwise, from the preferred alternative presented in the proposed plan. In addition, the State of South Carolina concurs with this remedy. South Carolina's letter of concurrence is provided in Appendix C to this ROD.

APPENDIX A - ANALYTICAL DATA SUMMARIES

Soil Analytical Data Summar Carolawn

Ft. Lawn, South Carolina

					1_01.7		1_QT.B	2-SLA	3 - 91. 1	3-SLB	4 –
STA		5-SLA	5-SLB	6-SLA	7-3LA		1-211	Z-5LA	3-SLA	3-216	4-
0211		5 5211	3 222				04/25/94	04/25/94	04/25/94	04/25/94	
04/2	25/94	04/25/94	04/25/94	04/2	26/94	04/	26/94				
INO	RGANIC	ELEMENTS			MG/KG		MG/KG	MG/KG	MG/KG	MG/KG	
MG/I	KG	MG/KG	MG/KG	MG/K	(G	MG/	KG				
	ARSE	_					1.2J	3.7	4.1	1J	3
			3	1.7J			0.0	0.5	<i>-</i> 1	400	
100	BARI	_	010	200	200		90	86	64	420	
1200		400 LLIUM	210			190		0.50J	0.44J	2.5JN	
0 7		.045J	1.5		 5J		4J	0.500	0.440	2.50N	
0.7.	CADM		1.5	0.75		0.7					
0.9	7J	_									
	COBA	LT			15		23	21	22	25	17
19		38	17	13							
		MIUM			62		38	83	39	40	380
170		-	210	220							
0.2.0	COPP		0.00	0.00	43			130			410
230	NICK		280	280	8.3		9.3	8.5J	8.4	46	21
22		.EL	35	14	0.3		9.3	0.50	0.4	40	21
22	LEAD		33		43		9.3	70	14		310
220		6.7	430	350							
	VANA	DIUM			51		100	77	110	130	63
50		73	67	53							
	ZINC				37J					58J	
1300	J 		43J	74J		120					
0 00	MERC	_		1 7				0.53			
0.98	3 7. T. T.T.M	1.0 IINUM		1.7	7900	0.7	13000	9400	16000	29000	
1200	OO ALUM		16000	1500)0	140		9400	16000	29000	
1200		ANESE	10000	1500	760		650	600	490	250	260
210	111110		550	180							
	CALC	!IUM			1800			1100	870	2600	

2600	2200	4700	2600	3300				
MAG	NESIUM		2000	1100	960	1200	15000	
3100	2700	7600	3000	3800				
IRO	N		18000	33000	22000	30000	42000	
25000	19000	27000	23000	20000				
SOD	IUM		120					
POT	ASSIUM		520	230	400	570	6000	550
780	1600	880	980					

* * * FOOTNOTES* * *

- ESTIMATED VALUE

- MATERIAL WAS ANALYZED FOR BUT NOT DETECTED

Appendix A-1

Soil Analytical Dat

UG/KG

Summary (cont)

Carolawn Ft. Lawn, South

UG/KG

Carolina							
			1-SLA		1-SLB		2-SLA
3-SLA	3-SLB	4-SLA		5-SLA		5-SLB	
6-SLA	7-SLA						
			04/25/94		04/25/94		04/25/94
04/25/94	04/25/94	04/25/94		04/25/94		04/25/94	
04/26/94	04/26/94						
PURGEABLE ORGANIC	COMPOUNDS		UG/KG		UG/KG		UG/KG
UG/KG	UG/KG	UG/KG		UG/KG		UG/KG	
UG/KG	UG/KG						
TRICHLOROETHE	ENE (TRICHLOROETHYLEN	E)					
							-
_	27Ј						
TETRACHLOROET	THENE (TETRACHLOROETH	YLENE)					
8J							
8J							
TOLUENE							
25J							
25J							
PESTICIDE/PCB COME	POUNDS		UG/KG		UG/KG		UG/KG

UG/KG

UG/KG

UG/KG

UG/KG	UG/KG					
4,4'-DDT 13	(P,P'-DDT) 				 	 _
4,4'-DDE 28	(P,P'-DDE) 				 	 -
PCB-1254 320	(AROCLOR 1254)	2900C	5400C	440	 	
PCB-1248	(AROCLOR 1248)		440C		 	 _
PCB-1260 	(AROCLOR 1260)		700C		 	 -

FOOTNOTES

J - ESTIMATED VALUE

- MATERIAL WAS ANALYZED FOR BUT NOT DETECTED

C - CONFIRMED BY GC/MS

Appendix A -2

Soil Analytical Data Summary (cont)

Carolawn

Ft. Lawn, South Carolina

			1-SLA	1-SLB	2-SLA
3-SLA	3-SLB	4-SLA	5-SLA	5-SLB	
6-SLA	7-SLA				
			04/25/94	04/25/94	
04/25/94	04/25/94	04/25/94	04/25/94	04/25/94	
04/25/94	04/26/94	04/26/94			
EXTRACTABLE OR	GANIC COMPOUNDS		UG/KG	UG/KG	
UG/KG	UG/KG	UG/KG	UG/KG	UG/KG	
UG/KG	UG/KG	UG/KG			
4 – N T	TROANILINE		190Ј		
					_
_					

	FLUORANTHENE		92J		
					-
-					
	PYRENE		110J		
					_
	CHRYSENE		180J		
					_
_					
	(3-AND/OR 4-)METHYLPHENOL				
		300J			
	PHENOXYBIPHENYL (2 ISOMERS)		900JN		
		-			_
_	 HEXACHLOROBIPHENYL (2 ISOMER:	g)	700JN		
			7000N 		_
_					
	(DIETHYLAMINO)PHENYLMETHANON	Ε	500JN		
					-
_	 AMINOANTHRACENEDIONE			80JN	
					_
_					
	DECAHYDROTRIMETHYLMETHYLENEM	ETHANO			
					_
_	 AZULENE				
2000JN					_
_					
	HEXAHYDROHYDROXYTRIMETHYL(ME	THYLETHYL)			
		-			_
_	PHENANTHRENONE (2 ISOMERS)			
1000JN		/ 			_
_					
	CEDROL				
1000JN					-
-	YLANGENE				
400JN					_
_					
	THUJOPSENE				
500JN		-			_
_	 QUATERPHENYL		300JN		
	·-	2000JN	4000JN	400JN	_
-					
	METHYLBENZOIC ACID				
700JN	 OXYBISBENZENE		700JN		
	OVIDIDDFINGFINE		/ U U U IN		

			2000JN		
600JN					
	CHLOROBIPHENYLOL				
		600JN			
	PHENOXYBIPHENYL				
		600JN			
	QUATERPRENYL (3 ISOMERS)				
	5 UNIDENTIFIED COMPOUNDS				
		10000J			_
_	10000J				
	9 UNIDENTIFIED COMPOUNDS				
8000JN					
		20000Ј			
	11 UNIDENTIFIED COMPOUNDS				
200000J					_
_					
	13 UNIDENTIFIED COMPOUNDS				
20000J				U0008	_
_					
	15 UNIDENTIFIED COMPOUNDS				
			70000J		_
_					
	17 UNIDENTIFIED COMPOUNDS				
					_
_	80000J				
	PETROLEUM PRODUCT				
		N		N	N

FOOTNOTES

J - ESTIMATED VALUE

N - PRESUMPTIVE EVIDENCE OF PRESENCE OF MATERIAL MATERIAL WAS ANALYZED FOR BUT NOT DETECTED

Appendix A-3

Soil Analytical Data Summary (cont)

Carolawn

Ft. Lawn, South Carolina

11-SLB		12-SLA	8-SLA	13-SLA	9-SLA	13-SLB			11-SLA
15-SLA 04/26/94 04/26/94	1	15-SLB 04/26/94 04/26/94	04/26/94			04/26/94			
MG/KG	IC ELEMENT	CS MG/KG MG/KG	MG/KG	MG/KG	MG/KG	MG/KG	MG/KG	MG/KG	MG/KG
	SILVER								
	ARSENIC		2.2						4.3
4.3				3.5				3.4	
2.4									
1.40	BARIUM		110		94	1.00	160		88
140 51		220 52		77		170		77	
21	BERYLLIU		0.84J		0.51J		0.92J		0.74J
	DERTHEE	0.88J	0.040	0.80J	0.510	0.89J	0.520	0.42J	0.740
0.35J				0.000		0.000		0.120	
	COBALT		13		12		21		7.7J
10		20		18		12		22	
10		9.5							
	CHROMIUM		54		32		33		33
9.3		47		69		41		75	
38	CORRER	40	0.3		2.0				4.5
33	COPPER	60	93	75	39	39		71	47
-				75		39		/ 1	_
	NICKEL		17		6.6		36		9.5J
9.3		29	_,	26		45		8.5	
6.2		12							
	LEAD		91		52		6.7		59
11J		92J		89J		2.9J		120Ј	
70J		33J							
	VANADIUM		65		59		69		58
69		72		75		63		72	
42	7 T N C	140	FOT						
37	ZINC	47	5ОЈ	38		43		36	
-				30		45		30	
	MERCURY		0.76						
		0.73						0.59	
0.32									
	ALUMINUM		15000		10000		15000		15000
23000		18000		14000		19000		13000	

8300	31000							
	MANGANESE	230		610		340		210
190	430		550		160		690	
160	160							
	CALCIUM	2800		1100		3600		1700
1400	5400		2900		5300		870	
1700								
	MAGNESIUM	4000		1700		8300		4600
6700	8700		5000		13000		900	
1400	1600							
	IRON	20000		18000		23000		20000
25000	25000		24000		24000		24000	
13000	50000							
	SODIUM							
								-
_	160							
	POTASSIUM	820		720		3000		1600
3000	4200		530		360		360	
E 4.0								
540	1000							

FOOTNOTES

J - ESTIMATED VALUE

-- - MATERIAL WAS ANALYZED FOR BUT NOT DETECTED

Appendix A-4

Soil Analytical Data Summar

(cont)

Carolawn Ft. Lawn, South

Carolina

11-SLB	12-SLA	13-SLA	13-SLB	14-SLA	8-SLA 15-SLA	9-SLA 15-SLB	10-SLA	11-SLA
04/26/94	04/26/94	04/26/94	04/26/94	04/26/94	4/26/94 04/26/	04/26/94 94 04/26/	04/26/94 94	04/26/94
PURGEABLE UG/KG	ORGANIC CO UG/KG	MPOUNDS UG/KG	UG/KG	UG/KG	UG/KG UG/KG	UG/KG UG/KG	UG/KG	UG/KG
TETR.	ACHLOROETHE 	NE (TETRACHL 	OROETHYLENE)	10J 			
PESTICIDE UG/KG	/PCB COMPOU UG/KG	NDS UG/KG	UG/KG	UG/KG	UG/KG UG/KG	UG/KG UG/KG	UG/KG	UG/KG
PCB-	1254 (AR	OCLOR 1254)			480			77
 ******	 *****	 *****	75 ******	28J ******	 ****			

FOOTNOTES

J - ESTIMATED VALUE

-- - MATERIAL WAS ANALYZED FOR BUT NOT DETECTED

Appendix A-5

Soil Analytical Data Summary (cont)

Carolawn

Ft. Lawn, South Carolina

GT 7	11 072	11 07 0	8-SLA	9-SLA	10-
SLA SLB	11-SLA 14-SLA	11-SLB 15-SLA	12-SLA 15-SLB	13-SLA	13-
STR	14-SLA	15-SLA	04/26/94	04/26/94	
04/26/	04/26/94	04/26/94	04/26/94	04/26/94	
04/26/		04/26/94	04/26/94	04/20/94	
04/20/	04/20/94	04/20/01	01/20/01		
EXTRAC	CTABLE ORGANIC COMPOUNDS		UG/KG	UG/KG	
UG/KG	UG/KG	UG/KG	UG/KG	UG/KG	
UG/KG	UG/KG	UG/KG	UG/KG		
F	BIS(2-ETHYLHEXYL) PHTHALAT	¬ г .			
M	METHYLIDYNEBENZENE		300JN		
					_
-					
Ç	QUATERPHENYL (3 ISOMERS)		1000JN		
					_
	 PHENYLTERPHENYL		300JN		
			3000N		
_					
Γ	DECAHYDROTRIMETHYLNETHANOA	AZULENE		800JN	
	==				_
_					
H	HEXAHYDROHYDROXYTRIMETHYL (METHYLETHYL)			
					_
_					
	PHENANTHRENOL (2 ISOMERS)				
900JN					_
_					
	HEXAHYDROHYDROXYTRIMETHYL (
					_
				300JN	
100JN	PHENANTHRENONE			3000N	_
T 0 0 0 IV					_
īv	METHYLHEXADIENE				
•					

300JI	N					-
-						
	PENTADECANOIC	ACID				
300JI	N					-
_	DITENTANTULDENO	(2 TGOMEDG)				
400JI	PHENANTHRENOL	(Z ISOMERS)				
-	LN					_
	METHYLBENZENE	SIILFONAMIDE				
			300JN			_
_						
	DICHLORONITRO.	ANILINE				
				70JN		-
-						
	DIISOCYANATOM	ETHYLBENZENE				
200JI						
	DECAHYDROTRIH	ETHYLMETHYLENEMETHA	NO			
						-
_	A CILL DATE					
	AZULENE					
 500JI	NT					
30001	COPAENE					
	COLLIENT					
			200JN			
	CHLORO (PHENYL	ENETHYL)PHENOL				
			200JN			
	OCTAHYDROTRIM	ETHYL(METHYLETHYL)				
						_
-	DITENTANTELLDENOT					
	PHENANTHRENOL					
			1000JN			
			100001			
	CHLORO (PHENYL	METHYL) PHENOL				
	•					_
_		70JN				
	OCTAHYDROTRIM	ETHYL(METHYLETHYL)P	HENANTHRENOL		2000JN	
						_
-		200JN				
	AMINOANTHRACE:	NEDIONE			800JN	
				200JN		-
-		100JN				
0.0 ===	HEXADECANOIC .	ACID				
80JN						-
_	1 11111111111111111111111111111		200JN	300JN	2000 T	
	1 UNIDENTIFIE	D COMBOOND			2000J	

			4000	 _
_		1000J		
2 UNID	ENTIFIED COMPOUNDS			
				 _
_	1000Ј			
4 UNID	ENTIFIED COMPOUNDS			
2000J		6000J		 _
-				
5 UNID	ENTIFIED COMPOUNDS			
3000J				
8 UNID	ENTIFIED COMPOUNDS			
				 _
-				
PETROL	EUM PRODUCT			
				 _
_				

FOOTNOTES

J - ESTIMATED VALUE

N - PRESUMPTIVE EVIDENCE OF PRESENCE OF MATERIAL

-- - MATERIAL WAS ANALYZED FOR BUT NOT DETECTED

Appendix A -6

Soi

Analytical Data Summary (cont)

Carolawn

Ft. Lawn, South Carolina

	16-SLA	17-SLA	18-SLA	19-SLA	
20-SLA	21-SLA	22-SLA	23-SLA	24-SLA	
25-SLA					

	04/26/94	04/26/94	04/26/94	04/27/94
04/27/94 04/26/94	04/27/94	04/26/94	04/26/94	04/26/94

INORGAN MG/KG MG/KG	NIC ELEMENT	'S MG/KG	MG/KG						
 1.9J	ARSENIC		1.9J	3.6		2.2J		1.8J	1.7J
50J 55	BARIUM	31J	43	120	43Ј	89	37J	38	52J

0.80J	BERYLLIUM	0.48J	0.42J			0.46J	0.49J	0.69J	0.58J
0.45J 8.2	COBALT	9.6	5.8	24	10	15	6.7	13	7.7
28	CHROMIUM	15	28	28	21	79	12	34	40
58 -	COPPER		40			81			
8.7 4.7J	NICKEL	6.8	3.8	12	9.5	3.9	13	4.1	12
9.8 18J	LEAD	4.8	45J	17	7.7	69Ј	6.3	13Ј	73
63 63	VANADIUM	29	59	65	32	39	22	48	60
 _	ZINC								24 -
14000 13000	ALUMINUM	9700	11000	14000	7400	7800	6300	9500	14000
89 500	MANGANESE	70	100	1100	130	740	130	360	120
 -	CALCIUM			1900	850J	1100	830J		
2500J 590	MAGNESIUM	1600J	1100	3300	2000Ј	640	2300Ј	480	2400Ј
24000 23000	IRON	11000	20000	21000	13000	12000	8700	16000	24000
1200 380	POTASSIUM	600	450	1300	630	320	300	280	840

FOOTNOTES

J - ESTIMATED VALUE

^{-- -} MATERIAL WAS ANALYZED FOR BUT NOT DETECTED

Soil Analytical Data Summary (cont Carolawn

Ft. Lawn, South Carolina

			16-SLA	17-SLA	18-SLA	19-SLA	20-SLA	21-SLA
22-SLA	23-SLA	24-SLA	25-SLA					
			04/26/94	04/26/94	04/26/94	04/27/94	04/27/94	04/27/94
04/26/94	04/26/94	04/26/94	04/26/9	4				
PURGEABLE	ORGANIC COM	IPOUNDS	UG/KG	UG/KG	UG/KG	UG/KG	UG/KG	UG/KG
UG/KG	UG/KG	UG/KG	UG/KG					
TOLUE	NE		5J		12			
		9Ј						

PESTICIDE/PCB COMPOUNDS NONE DETECTED

* * * FOOTNOTES* * *

J - ESTIMATED VALUE

-- - MATERIAL WAS ANALYZED FOR BUT NOT DETECTED

Appendix A-8

Soil Analytical Data Summary (cont)

Carolawn

Ft. Lawn, South Carolina

			16-SLA	17-SLA	18-SLA
19-SLA	20-SLA	21-SLA	22-SLA	23-SLA	
24-SLA	25-SLA				
			04/26/94	04/26/94	
04/26/94	04/27/94	04/27/94	04/27/94	04/26/94	
04/26/94	04/26/94	04/26/94			
EXTRACTA	BLE ORGANIC COMPOUNDS		UG/KG	UG/KG	UG/KG
UG/KG	UG/KG	UG/KG	UG/KG	UG/KG	
UG/KG	UG/KG				
DII	SOCYANATOMETHYLBENZENE		200JN		
					_
_					
DEC	AHYDROTRIMETHYLMETHANOAZULENE			3000JN	200JN
					-
-					
(DI	NETHYLETHYL)METHYLPHENOL				
1000JN					
_					
(HY	DROXYPHENYL) ETHANONE				

			100JN			-
_						
DECAHYDROTR	IMETHYLMETHYLENE					
_						_
METHANOA	ZULENE					
			100JN			_
_						
(HYDROXYMET	HYL) ETHANONE					
			100JN			-
-						
PHENANTH	RENOL			500	0JN	
1000JN			400JN			-
-		1371 DM11371 \				
HEXAHYDROHY	DRODXYTRIMETHYL(METE		 			
_						
DECAHYDROTR	IMETHYLMETHYLENEMETH	IANO				
						-
-						
AZULENE						
	200JN	900JN				
COPAENE						
200.TN						
200JN HEXADECANOT	 C ACID					
200JN HEXADECANOI	C ACID		 200JN		400JN	
HEXADECANOI	C ACID 		 200JN		400JN	
HEXADECANOI 200JN	C ACID IMETHYL(METHYLETHYL)	 PNENANTHRENE	 200JN		400JN	
HEXADECANOI 200JN	 	 PNENANTHRENE 	 200JN		400JN	
HEXADECANOI 200JN	 		 200JN		400JN	
HEXADECANOI 200JN	 	 PNENANTHRENE 100JN	 200JN		400JN	
HEXADECANOI 200JN OCTAHYDROTR	 IMETHYL (METHYLETHYL) 	 100JN	 200JN		400JN	
HEXADECANOI 200JN OCTAHYDROTR OCTAHYDROTR	 	 100JN PHENANTHRENOL	 200JN		400JN	
HEXADECANOI 200JN OCTAHYDROTR	 IMETHYL (METHYLETHYL) 	 100JN	 200JN		400JN	
HEXADECANOI 200JN OCTAHYDROTR OCTAHYDROTR	 IMETHYL (METHYLETHYL) 	 100JN PHENANTHRENOL 300JN	 200JN		400JN	
HEXADECANOI 200JN OCTAHYDROTR OCTAHYDROTR	 IMETHYL (METHYLETHYL) 	 100JN PHENANTHRENOL 300JN	 200JN		400JN	
HEXADECANOI 200JN OCTAHYDROTR OCTAHYDROTR	IMETHYL (METHYLETHYL) IMETHYL (METHYLETHYL)	 100JN PHENANTHRENOL 300JN	 200JN		400JN	
HEXADECANOI 200JN OCTAHYDROTR OCTAHYDROTR 1000JN	IMETHYL (METHYLETHYL) IMETHYL (METHYLETHYL)	 100JN PHENANTHRENOL 300JN	 200JN		400JN	
HEXADECANOI 200JN OCTAHYDROTR OCTAHYDROTR 1000JN	IMETHYL (METHYLETHYL) IMETHYL (METHYLETHYL)	100JN PHENANTHRENOL 300JN	 200JN		400JN	
HEXADECANOI 200JN OCTAHYDROTR OCTAHYDROTR 1000JN	IMETHYL (METHYLETHYL) IMETHYL (METHYLETHYL)	 100JN PHENANTHRENOL 300JN	 200JN		400JN	
HEXADECANOI 200JN OCTAHYDROTR OCTAHYDROTR 1000JN (2 ISOME	IMETHYL(METHYLETHYL) IMETHYL(METHYLETHYL)	100JN PHENANTHRENOL 300JN 2000JN	 200JN		400JN	
HEXADECANOI 200JN OCTAHYDROTR OCTAHYDROTR 1000JN (2 ISOME DECAHYDROTR	IMETHYL (METHYLETHYL) IMETHYL (METHYLETHYL)	100JN PHENANTHRENOL 300JN 2000JN	 200JN		400JN	
HEXADECANOI 200JN OCTAHYDROTR OCTAHYDROTR 1000JN (2 ISOME	IMETHYL(METHYLETHYL) IMETHYL(METHYLETHYL)	100JN PHENANTHRENOL 300JN 2000JN	 200JN		400JN	
HEXADECANOI 200JN OCTAHYDROTR OCTAHYDROTR 1000JN (2 ISOME DECAHYDROTR	IMETHYL(METHYLETHYL) IMETHYL(METHYLETHYL)	100JN PHENANTHRENOL 300JN 2000JN	 200JN		400JN	
HEXADECANOI 200JN OCTAHYDROTR OCTAHYDROTR 1000JN (2 ISOME DECAHYDROTR	IMETHYL(METHYLETHYL) IMETHYL(METHYLETHYL)	100JN PHENANTHRENOL 300JN 2000JN	 200JN		400JN	
HEXADECANOI 200JN OCTAHYDROTR OCTAHYDROTR 1000JN (2 ISOME DECAHYDROTR 700JN	IMETHYL(METHYLETHYL) IMETHYL(METHYLETHYL)	100JN PHENANTHRENOL 300JN 2000JN	 200JN		400JN	
HEXADECANOI 200JN OCTAHYDROTR OCTAHYDROTR 1000JN (2 ISOME DECAHYDROTR	IMETHYL(METHYLETHYL) IMETHYL(METHYLETHYL)	100JN PHENANTHRENOL 300JN 2000JN	 200JN		400JN	
HEXADECANOI 200JN OCTAHYDROTR OCTAHYDROTR 1000JN (2 ISOME DECAHYDROTR 700JN THUJOPSENE	IMETHYL(METHYLETHYL) IMETHYL(METHYLETHYL)	100JN PHENANTHRENOL 300JN 2000JN	200JN		400JN	

3000	JN							
	OCTADECANOIC	ACID						300JN
								-
-		300JN						
	OCTAHYDROTRI	METHYL(METHYLETHYL)						
								-
-								
	PHENANTHR	ENOL (2 ISONERS)						
		3000JN	3000JN					
	\							
5000			, D. I.D.					
	METHYL (TRIME	THYLCYCLOPENTYL)BENZ 300JN	ENE					
		3000N						
3000	TNI							
3000		ETHYL(METHYLETHYL)PH	IENANTHRENE					
	OCTAILI DRODIN							_
_								
	CARBOXYLI	C ACID, METHYLESTER						
4000	ΙΝ							
	HEXAHYDROHYD	ROXYTRIMETHYL(METHYI	ETHYL)					
								-
-								
	PHENANTHR							
		700JN						
		300JN						
		300JN	200JN					
3000		ED COMPOUNDS						
	Z UNIDENIIFI	ED COMPOUNDS			1000Ј			
					10000			
	3 UNIDENTIFI	ED COMPOUNDS		100	00J			2000J
	3 011111111							20000
1000)J							
		ED COMPOUNDS				300	0J	
								_
_								
	S UNIDENTIFI	ED COMPOUNDS						
		7000J					5000J	-
-								
	6 UNIDENTIFI	ED COMPOUNDS						
5000								
0000		ED COMPOUNDS						
8000	JU							_
_								

11 UNIDENTIFIED COMPOUNDS -- -- -- 10000J -- -- --

FOOTNOTES

J - ESTIMATED VALUE

N - PRESUMPTIVE EVIDENCE OF PRESENCE OF MATERIAL

-- - MATERIAL WAS ANALYZED FOR BUT NOT DETECTED

Appendix A-9

Soi

Analytical Data Summary (cont)

Carolawn

Ft. Lawn, South Carolina

			26-SLA				27-SLA		28-SLA
29-S		30-SLA		31-SLA		32-SLA		33-SLA	
33-S	LB	•	4.400.404		4.407.404	•	4.400.404		4 / 0 7 / 0 4
04/0	C / O A		4/27/94 94		4/27/94	0 / 04/27	4/27/94		4/27/94
	6/94 7/94	04/2//	94	04/2//	94	04/2//	94	04/27/	94
04/2	7/94								
INOR	GANIC ELEMENTS		MG/KG		MG/KG		MG/KG		MG/KG
MG/K	G	MG/KG		MG/KG		MG/KG		MG/KG	
MG/K	G								
	ARSENIC		2.6		4				2 J
		2.8		1.9J		2J		2.5	-
_			5 2-		000-		10-		0.4.=
35	BARIUM	C C T	53J	ГОТ	200J	267	40J	ГЛТ	24J
35 100J		66J		50J		36J		54J	
1000	BERYLLIUM		0.77J		4.1JN		0.41J		O.44J
0.38		1.2	0.770	0.61J	1.101	0.48J	0.110	0.62J	0.110
1J									
	COBALT		8.1		28		3.5		2.1
8		8.6		3.7		2.9		5.2	
14									
	CHROMIUM		25		16		16		13
21		21		17		14		28	
93	COPPER								17
	COPPER								
_									
	NICXEL		7.3		18		4.8		3.6
		8.3		4.7		2.4J		6.6	

56									
9.8J	LEAD	15	20	12		13	14	11	6.5
6.7									
	VANADIUM	100	69		110	5 0	45	0.5	41
44 73		100		44		53		97	
7 3	ZINC				65				
9.6						15			
53	ALUMINUM		16000		16000		14000		9000
7600		23000	10000	10000	10000	10000	14000	20000	9000
2500									
	MANGANESE		82		300		56		18
210 140		84		64		30		70	
110	CALCIUM		940J		5600J				
				370J					_
-	NAN CALL CITINA		1600J		9200Ј		0007		2707
620	MAGNESIUM	1700J	10000	480J	92000	420J	920J	1400J	370Ј
1100	0J								
	211011		25000		41000		17000		14000
1400 3400		35000		17000		18000		35000	
	POTASSIUM		790		5200		710		240
310		940		410		240		820	
1500									

FOOTNOTES

J - ESTIMATED VALUE

N - PRESUMPTIVE EVIDENCE OF PRESENCE OF MATERIAL

-- - MATERIAL WAS ANALYZED FOR BUT NOT DETECTED

pendix A -10

Soil Analytical Data Summary (cont)

Carolawn

Ft. Lawn, South Carolina

			26-SLA	26-SLB	27-
SLA	28-SLA	29-SLA	30-SLA	31-SLA	32-
SLA	33-SLA	33-SLB			
			04/27/94	04/27/94	
04/27/94	04/27/94	04/26/94	04/27/94	04/27/94	

Аp

04/27/94	04/27/94	04/27/94			
EXTRACTABLE ORG	ANTC COMPOUNDS		UG/KG	UG/KG	
UG/KG	UG/KG	UG/KG	UG/KG	UG/KG	
UG/KG	UG/KG	UG/KG	OG/ NG	0 G / RG	
UG/KG	UG/ KG	UG/KG			
OCTADECANO	DIC ACID			300JN	
					_
_					
OCTAHYDROD	IMETHYL (METHYLETHYL)PI	HENANTHRENE			
					_
_					
CARBOXY	LIC ACID, METHYLESTER			300JN	
					_
_					
OCTAHYDROD	IMETHYL(METHYLETHYL)				
					_
_					
PHENANT	HRENECARBOXYLIC ACID,	METHYLESTER			
300JN					_
_					
OCTAHYDROT	RIMETHYL (METHYLETHYL)	PHENANTHRENOL			
					_
_					
(2 ISOM	IERS)				
4000JN					_
_					
OCTAHYDROM	IETHYLMETHYLENE (METHYL)	ETHVI.)			
					_
_					
METHANC	TNDENE				
	600JN				_
_					
DECZHADBUL	RIMETHYLMETHYLENEMETH	ANOAZIII.ENE			
	3000JN				_
_	30000N				
	DIMETMYL(METHYLETHYL)PI	υΓΝΛΜͲϤΟΓΝΓ			
OCIAIIIDROL					_
_					
CAPROYY	LXC ACID, METHYLESTER				
CARBOXI	1000JN				
	100001				_
-					
YLANGENE		200 TM			
		200JN			_
		A DIIMIIA T HATONIH			
OCTAHYDROT	'ETRAMETHYLCYCLOPROPANA				
		300JN			_
-				400	
CEDROL			400 777	400JN	
			400JN		_
- -	 ,				
THUJOPSENE		0.0.0.	400===		
	900JN	200JN	400JN		_

	METHYL (TRIMETHYLCYCLOPENT	YL)BENZENE			
	, NC008	, 	500JN		_
_					
	HEXAHYDROTRIMETHYL (METHYL)	ETHYL)			
					-
_					
	PHENANTHRENONE				
			400JN		-
_				4000 777	
	PHENANTHRENOL	2000 TM		4000JN 	
 200J		3000JN	7000JN		
2000	DECAHYDROTRIMETHYLMETHANO	N ZIII. ENIE			
	900JN				
		500JN			
1000					
900J					
900J					
	HEXADECANOIC ACID				
	400JN				
700J					
600J	N	600JN			
	OCTADECENOIC ACID				
800J		N0008			
	OCTAHYDROTRIMETHYL (METHYL)	ETHYL)			
					-
-		>			
	PHENANTHRENOL (2 ISOME				
		3000JN			
	6000JN 				
4000					
4000					
4000	OCTAHYDRODIMETHYL (METHYLE'	THYI.) DHENANTHRENE			
					_
_					
	CARBOXYLIC ACID, METHY	LESTER (2 ISOMERS)			
1000	JN	1000JN			
	1 UNIDENTIFIED COMPOUND				
		200J			
		2000Ј			
	2 UNIDENTIFIED COMPOUNDS		2000Ј		
					_
					
	3 UNIDENTIFIED COMPOUNDS				

		3000JN			_
-					
	4 UNIDENTIFIED COMPOUNDS				
6000J					_
-					
	7 UNIDENTIFIED COMPOUNDS			4000J	
					_
-	4000J				
9	9 UNIDENTIFIED COMPOUNDS				
			7000J		_
-					
:	12 UNIDENTIFIED COMPOUNDS				
10000	J				_
-					

FOOTNOTES

- ESTIMATED VALUE J

- PRESUMPTIVE EVIDENCE OF PRESENCE OF MATERIAL
- MATERIAL WAS ANALYZED FOR THE PROPERTY OF T - MATERIAL WAS ANALYZED FOR BUT NOT DETECTED

Appendix A -11

Soi

Analytical Data Summary (cont)

Carolawn

Ft. Lawn, South Carolina

		26-SLA	26-SLB	27-SLA	28-SLA
29-SLA	30-SLA	31-SLA	32-SLA	33-SLA	
33-SLB					
		04/27/94	04/27/94	04/27/94	
04/27/94	04/26/94	04/27/94	04/27/94	04/27/94	
04/27/94	04/27/94				
PURGEABLE ORGAN	IIC COMPOUNDS	UG/KG	UG/KG	UG/KG	UG/KG
UG/KG	UG/KG	UG/KG	UG/KG	UG/KG	
UG/KG					
TOLUENE					
	2J				-
_					

PESTICIDE/PCB COMPOUNDS NONE DETECTED

FOOTNOTES

J - ESTIMATED VALUE

-- - MATERIAL WAS ANALYZED FOR BUT NOT DETECTED

Appendix A -12

STRONTIUM

TITANIUM

NA

47

NA

1200

NA

NA

NA

34

NA

720

Anal	vtical Data S	ummary (cont)				Soi
	,					
Caro	lawn					
	a .1 a	7.				F
t. L	awn, South Ca	rolina				
		34-SLA	35-SLA	35-SLB	36-SLA	
37-S	LA	38-SLA	39-SLA	40-SLA	41-SLA	
45-S		45-SLB				
		04/27/94	04/27/94	04/27/94	04/26/94	
04/2	6/94	04/26/94	04/26/94	04/26/94	04/26/94	
10/2	5/94	10/25/94				
	GANIC ELEMENT		MG/KG	MG/KG	MG/KG	
MG/K		MG/KG	MG/KG	MG/KG	MG/KG	
MG/K	.G	MG/KG				
	ARSENIC	2.6			1.8J	
5.7		2.3J	2.9	2.9	23	_
-						
	BARIUM	52J	40J	36	51	
72		130	98	85	140	
100		58				
	BERYLLIUM		0.55J		0.42J	
0.51	J	0.89J	1.1J	0.49J		-
-						
	COBALT	13	4.9	3.2	8.8	
12		18	22	8.9	7.3	
12		5.3				
	CHROMIUM	14	62	15	32	
38		34	85	33	220	
14		42				
	COPPER	20		20	15	
		49	30		68	
30		39				
	NICKEL	2J		3.5	3.3J	
8.9		26	14	7	8.7	
11		20				
	LEAD	15	9.7	8	15J	
14J		14J	17J	9.7J	280J	
22		7.7				

NA

F

VANADIUM	46	47	72	45
49	75	120	57	31
76	34			
YTTRIUM				
13	11			
ZINC				
				110
38	34			
MERCURY				
0.32	0.17			
ALUMINUM	9300	7800	25000	9200
8600	17000	14000	10000	10000
20000	14000			
MANGANESE	120	66	40	410
430	360	590	270	230
230	120			
CALCIUM				800
1900	3300	1500	1400	49000
2900	830			
MAGNESIUM	510J		1100J	520
2100	6500	2600	2500	26000
4800	3000			
IRON	17000	20000	29000	15000
15000	24000	34000	17000	16000
25000	15000			
POTASSIUM	400	250	830	310
780	1800	750	920	2200
2400	2300			
SODIUM				
_	200			

* * *FOOTNOTES* * *

J - ESTIMATED VALUE

-- - MATERIAL WAS ANALYZED FOR BUT NOT DETECTED

NA - NOT ANALYZED

Appendix A -13

Soil Analytical Data Summary (cont)

Carolawn

Ft. Lawn, South Carolina

			34-SLA	35-SLA
35-SLB	36-SLA	37-SLA	38-SLA	39-SLA
40-SLA	41-SLA	45-SLA	45-SLB	

04/27 04/26		04/26/94 10/25/94	04/27/94 04/26/94 10/24/94	04/27/94 04/26/94	
EXTRA	CTABLE ORGANIC COMPOUNDS		UG/KG	UG/KG	
UG/KG		UG/KG	UG/KG	UG/kG	
UG/KG		UG/KG	UG/KG		
	4-NITROANILINE	, -			
					_
_	3800J				
	OCTAHYDRODIMETHYL (METHYLETHYL) PI	HENANTHRENE			
					_
-					
	CARBOXYLIC ACID, METHYLESTER	(2 ISOMERS)	600JN		
					_
_					
	DECAHYDROTRIMETHYLMETHANOAZULEN	E		500JN	
					_
_					
	METHYL (TRIMETHYLCYCLOPENTYL)BEN	ZENE		100JN	
					-
-					
	HEXADECANOIC ACID				
500JN	- -				_
-					
	HEXAHYDRODIMETHYL (METHYLETHYL)				
					_
_	 NAPHTHALENE				
	NAPHIHALENE 100JN				
	1000N				_
_	OCTAHYDROTRIMETHYL(METHYLETHYL)				
	OCIAIIDKOIKIMEIIIID(MEIIIIDEIIIID)				_
_					
	HEXAHYDROXYTRIMETHYL(METHYLETHY	L) PHENANTHRENONE			
	100JN				_
_					
	(DIMETHYLETHYL)PHENOL				
		500JN			_
_					
	DECAHYDROTRIMETHYLMETHYLENEMETH	ANOAZULENE			
			300JN		_
_					
	OCTAHYDROTRIMETHYL (METHYLETHYL)				
					_
-					
	PHENANTHRENOL (2 ISOMERS)				
	700JN		5000JN		_
-					
	METHYLTRIMETHYLCYCLOPENTYLBENZE	NE			
				200JN	-
-					
	METHYLPHENYLANTHRACENEDIONE			1000	
				1000JN	-

-					
I	HEXAHYDROHYDROXYTRIMETHYLMET	HYLETHYL			
					-
-					
(OCTAHYDROTRIMETHYL (METHYLETH	YL)PHENANTHRENOL	300JN		
				7000JN	_
_					
ī	DECAHYDROTRIMETHYLMETHYLENEM:	ETHANO			
					_
_					
	א וקוון האום				
	AZULENE			1000 TN	
				1000JN	
2000JI					
(CEDROL				
400JN					
-	THUJOPSENE				
300JN					
1	METHYL(TRIMETHYLCYCLOPENTYL)	METHYLBENZENE			
300JN					
	OCTAHYDROTRIMETHYL(METHYLETH	VI \ DUENIANTUDENONE			
(JCIAHIDROIRIMEIHIL (MEIHILEIH	IL / PHENANTHRENONE			
					_
-				0.0.0.	
	(2 ISOMERS)			2000JN	
					
7000JI	N				
I	HEXAHYDROHYDROXYTRIMETHYL(ME	$\mathtt{THYLETHYL}$)			
					_
-					
	PHENANTHRENONE				
			1000JN	800JN	
600JN					
	(DIMETHYLETHYL)METHYLPHENOL				
300JN					_
3000N	10000JN				
1	ANTHRACENEDIONE				
					_
-	6000JN				
-	1 UNIDENTIFIED COMPOUND				
		1000Ј			-
-					
2	2 UNIDENTIFIED CONPOUNDS			300J	
	4000J				_
_					
ļ	5 UNIDENTIFIED COMPOUNDS		8000J		
					_
_					
,	6 UNIDENTIFIED COMPOUNDS				
10000			· -		
T0000	J				_
_					
_	 8 UNIDENTIFIED COMPOUNDS				

5000J 700000J --*****************

FOOTNOTES

J - ESTIMATED VALUE
N - PRESUMPTIVE EVIDENCE OF PRESENCE OF

MATERIAL

-- - MATERIAL WAS ANALYZED FOR BUT NOT DETECTED

Appendix A -14

Soil Analytical Data Summary (cont)

Carolawn

Ft. Lawn, South Carolina

			34-SLA	35-SLA	35-SLB
36-SLA	37-SLA	38-SLA	39-SLA	40-5	SLA
41-SLA	45-SLA	45-SLE	3		
			04/27/94	04/27/94	04/27/94
04/26/94	04/26/94	04126/94	04/26/9	94 04/2	26/94
04/26/94	10/25/94	10/25/	94		
PURGEABLE ORGAN	IC COMPOUNDS		UG/KG	UG/KG	UG/KG
UG/KG	UG/KG	UG/KG	UG/KG	UG/E	(G
UG/KG	UG/KG	UG/KG			
TETRACMLOR	DETHENE (TETRACHLORO	ETHYLENE)	3Ј	5J	
TOLUENE			6J	11J	
PESTICIDE/PCB CO			UG/KG	UG/KG	UG/KG
UG/KG	UG/KG	UG/KG	UG/KG	UG/E	KG .
UG/KG	UG/KG	UG/KG			
PCB-1254 (2	AROCLOR 1254)				
780					
4,4'-DDE					
15J					
15J					
*****	* * * * * * * * * * * * * * * * * * * *	******	******	k *	

FOOTNOTES

- J ESTIMATED VALUE -- MATERIAL WAS ANALYZED FOR BUT NOT DETECTED

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		Analytical Data Sun Carolawn Ft. Lawn, South Car	
3-SD	4-SD	1-SD	2-SD
04/26/94	04/26/94	04/26/94	04/26/94
INORGANIC ELEMENT		MG/KG	MG/KG
MG/KG	MG/KG		
ARSENIC			
BARIUM 24			
BERYLLIUM -		0.30J	
COBALT	3.7	1.4J	3.1
CHROMIUM	6	7.1	11
LEAD 2.1J	1.8	1.8	2.1J
VANADIUM 18	12	15	13
ZINC	14		
- ALUMINUM		1100	1600
1500 MANGANESE	1100	270	290
310 IRON	250	6200	5100
7800 POTASSIUM	5500	76	88
62	140		
PURGEABLE ORGANIC PESTICIDE/PCB COM		NONE DETECTED NONE DETECTED	
EXTRACTABLE ORGAN	IC COMPOUNDS UG/KG	UG/KG	UG/KG
	OG/ RG HYLMETHYLENE (METHYLETHYL)NAPHTHALENE		
-	600JN		_
OCTADECENOIC	ACID		

-	900JN		
	OCTADECANOIC ACID	 	-
-	900JN		
	OCTAHYDROTRIMETHYL (METHYLETHYL)PHENANTHRENOL	 	_
-			
	(2 ISOMERS)	 	_
_	8000JN		
	15 UNIDENTIFIED COMPOUNDS	 	-
-	20000J		
	1 UNIDENTIFIED COMPOUND	 900Ј	_
-			

FOOTNOTES

J - ESTIMATED VALUE -- - MATERIAL WAS ANALYZED FOR BUT NOT DETECTED

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Surface Wa=er Analytical Data Summary

Carolawn

- Ft. Lawn, Sou=h Carollna

		1-SW	2-SW	3-SW	4-SW
201-SW	401 -TB	04/26/94	04/26/94	04/26/94	
04/26/94	04/26/94	04/25/94			
INORGANIC ELEMENTS		UG/L	UG/L	UG/L	UG/L
UG/L	UG/L				
BARIUM		33	29	32	30
32 STRONTIUM	NA	95	86	93	89
93 TITANIUM	NA	7.3	6.2	6.2	5.8
7.4	NA				
ALUMINUM 310	NA	350	280	260	240
MANGANESE 59	NA	60	52	53	59
	1411				
MG/L	MG/L	MG/L	MG/L	MG/L	MG/L
CALCIUM		9.3	8.4	9.1	8.6
9.1	NA				
MAGNESIUM 3.8	NA	3.8	3.5	3.8	3.6
IRON	NIA	0.88	0.76	0.80	0.69
0.85	NA				

п. о	SODIUM	7.9	7.2	7.9	7.6	
7.8	POTASSIUM	NA	1.5	1.4	1.6	1.5
1.6		NA				

PURGEABLE ORGANIC COMPOUNDS NONE DETECTED PESTICIDE/PCB COMPOUNDS NONE DETECTED EXTRACTABLE ORGANIC CONFOUNDS NONE DETECTED

FOOTNOTES

NA = NOT ANALYZED

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Field Parameter Data Summar Carolawn

Ft. Lawn, South Carolina

Sample Number	pH (SU)	Specific Conductance (umhos/cm@25øC)	Temperature (øc)
001-SW	6.9	202	19.5
002-SW	6.3	122	20.0
003-SW	6.3	121	21.6
004-SW	6.7	121	23.8

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APPENDIX B - RESPONSIVENESS SUMMAR

RESPONSIVENESS SUMMAR CAROLAWN (OU2) SUPERFUND SITE

1. Overview

The U. S. Environmental Protection Agency (EPA) held a public comment period from July 24, 1995 to August 24, 1995, for interested parties to comment on the Remedial Investigations and the Baseline Risk Assessment results and the Proposed Plan for the Carolawn (OU2) Superfund Site in Fort Lawn, South Carolina. The comment period closed on August 24, 1995.

EPA held a public meeting at 7:00 p.m. on August 10, 1995 at the Lewisville Elementary School in Fort Lawn, South Carolina to

present the results of the Remedial Investigation and the Baseline Risk Assessment, to present the Proposed Plan and to receive comments from the public.

In the absence of any significant source of contamination in the soil, surface water and sediment at the Site, the No Action alternative was proposed by EPA to address the soil, surface water and sediment. In addition, a groundwater remedy has been selected under a Record of Decision for Carolawn (OUI). However, should future monitoring of the site (e.g. Five-Year Review) indicate that the site poses an unacceptable risk to the environnment, then EPA, in consultation with the State of South Carolina, may initiate clean-up actions under the authority of CERCLA nd in accordance with the National Oil and Hazardous substances Pollution Contingency Plan. Judging from the comments received during the public comment period, the residents and local officials in the Fort Lawn, South Carolina area support the cleanup alternative proposed by EPA.

The Responsiveness Summary provides a summary of citizens' comments and concerns identified and received during the public comment period, and EPA's response to those comments and concerns. These sections and attachments follow:

Background of Community Involvement

Summary of Comments Received During the Public Comment Period and EPA's Responses

Attachment A: Proposed Plan for the Carolawn (OU2) Superfund Site

Attachment B: Public Notices of Public Comment Period

Attachment C: Written Public Comments Received During the Public Comment Period

Attachment D: Official Transcript of the Proposed Plan Public Meeting

2. Background of Ccmununity Involvement

EPA's community relations program for the Site began in 1987, when EPA conducted community interviews in order to develop a community relations plan for the Site. At that time, residents living adjacent to the Site were concerned about the Site and about any health risks from the Site. In addition, residents did voice some concerns about lack of information to the public during the removal work at the Site and lack of response to earlier complaints about the Site.

Throughout EPA's involvement, the community has been kept aware and informed of Site activities and findings. Discussions have taken place during visits to the area by the Remedial Project Manager (RPM) and the Community Relations Coordinator (CRC). Concerned citizens and Local officials were briefed prior to the Proposed Plan Public Meeting held on August 10, 1995 The Site mailing list was expanded to include additional residents living in close proximity to the Site.

3. Summary of Comments Received During the Public Comment Period and Agency Responses

The Public Comment Period was opened on July 24, 1995 and was ended on August 24, 1995. Public Notices which were published in local papers can be found in Attachment B.

On August 10, 1995, EPA held a public meeting to present the Proposed Plan to the community and to receive comments thereupon. All comments received at this public meeting and during the public comment period are summarized below.

Summary and Response to Local Community Concerns

The following issues and concerns were expressed at the Proposed Plan Public Meeting, and during the public comment period.

COMMENT: Several citizens expressed a concern that the Remedial Investigation did not encompass the entire parcel of property of 60 acres and would like an additional investigation to take place on the adjacent acreage. Moreover, several citizens are apprehensive about the existence of buried drums and feel that an additional investigation would alleviate their concerns.

RESPONSE: Previous studies suggested that there were numerous sources of contamination at the Site. Based on those studies, several remedial actions have been performed to remove contaminated soils, drums (some buried) and liquid waste frou the Site. While those levels of contamination were greatly reduced, several Remedial Investigations were warranted to fully delineate all contamination of known areas and to characterize the Site. Based on the information obtained from the operational history of the facility and the earlier investigations, including this Remedial Investigation, EPA has characterized the Site and the nature of its contaminants at all known areas of contamination. However, if further information (i.e., via the Citizen Advisory Group) suggests additional sources of contamination exist, EPA will investigate the area of concern to confirm the nature and extent of contamination on. any of the remaining acreage.

COMME: An attendee requested EPA to appoint a committee from the community to participate with the agency in future efforts and

decisions for the Site.

RESPONSE: Based on citizen interest at the meeting, EPA will pursue the establishment of a Citizen Advisory Group for the Carolawn Site. Once this group is established, the Citizen Advisory Group will participate with EPA in future efforts and decisions for the Site. In addition, formation of the Citizens Acvisory Group will increase dissemination of information and provide viable feedback from the community for on-going implementation issues as well as determining the need for additional investigation on the remaining acreage.

COMMENT: An attendee expressed a concern that the Carolawn Site was cited as one of 114 sites in the United States that most needed cleaning up.

RESPONSE: Upon completion of operational practices which occurred during the 1970's, the Carolawn property was an area covered with two incinerators, several storage tanks, two storage trailers and many drums (both inside and outside the 3-acre fenced area). During the early 1980 's, SCDHEC and EPA conducted several site investigations at the Carolawn Site. The results of these investigations showed the presence of trichlorethane (TCE) and other solvents in nearby residential wells. The results also indicated that the Site was contaminated with big levels of metals and organic compounds.

During the late 1970's and early 1980's, the Site could have been perceived as one of the worst sites in the United States. However, due to the elevated levels of contamination found and the potential threat for imminent damage to public health and/or the environment, EPA initiated cleanup activities at the Site on December 1, 1981. The cleanup activities continued through February 1982, and included removal of contaminated soils, drums (some buried), and

liquid wastes from the Site. Due to each of the response action that have occurred at the Site, the levels of contamination have been greatly reduced.

Currently, the Site does not pose an imminent threat to public health and/or the environment. However, the Site does pose a long-term threat to the public health through exposure to the groundwater. A remedy has been selected for groundwater remediation at the Site and is expected to be implemented in the near future.

COMMENT: An attendee inquired about whether or not there is additional funding to support any further testing of the other 60 acres of the Site.

RESPONSE: In response, EPA stated that the Agency's current status for funding is questionable. Based on budget cuts and the

occurrence of a Recision Bill that was passed this year to basically pull back funds allocated for 1995, Region IV has shut down some starts of some sites in other states that were ready to implement cleanup activities. As far as we know, EPA has funds for next year. However, the Agency does not know how long the Superfund program will have funds. Like other Federal agencies, funding for EPA has to be appropriated each year. Unfortunately, the Superfund Law does not expire, but the part of the Law that collects the tax that generates the money to fund the program does expire. Thus, the program could go on if there is funding in the trust fund to continue on. At this time, the agency is not sure about

~

reauthorization or when the Superfund Law will be reauthorized. Therefore, it is hard to commit to saying there will be funding for the kinds of investigations we would have to do. Currently, EPA will have to start prioritizing everything to the worst-case-first scenario. That being the case, further investigation of this Site might not break out as a worst-case-scenario if there are limited funds. EPA will try to obtain additional funds and continue to go forward and maybe even do some things in-house of a limited nature with the existing resources in-house. If the Agency has solid leads, we could also work through SCDHEC to try to pursue things

that way. At this point, it is an unanswerable question but, there are options available. We think the Agency willhave funds, and we think that if there is a legitimate need, the Agency will go forward and investigate those things.

Attachment A

Proposed Plan for the Carolawn (OU2) Superfund Site

SUPERFUND PROPOSED PLAN FACT SHEET
Carolawn Superfund Site-Operable Unit Two
Fort Lawn, Chester County, South Carolina

U.S. Environmental Protection Agency, Region IV, Atlanta, GA July 1995

This fact sheet is one in a series designed to inform comment period has ended residents and local officials of the ongoing cleanup efforts to EPA during that time has at the Site. A number of terms specific to the Superfund

the Site only after the public and all information submitted been reviewed and considered.

As outlined in section process (printed in bold print) are defined in the glossary 117(a) of CERCLA, EPA encourages public participation which begins on Page 12. for addressing contamination

providing an opportunity for the INTRODUCTION proposed remedial actions.

alternative, or a change from the The United States Environmental Protection Agency (EPA) another, may be made if public presents this Proposed Plan for no further action for the indicate that such a change Carolawn (OU2) Superfund Site ("the Site"), located in Fort would result in a more appropriate solution. The final Lawn, Chester County, South Carolina. Contaminant levels remedy will be documented have been substantially reduced through implementation of after EPA has taken into soil and source area cleanup activities conducted through a from the public. Upon timely Removal Action which occurred December 1981 through public comment period by 30 February 1982. In addition, a groundwater remedy has been selected for Carolawn (OU1). Studies to date indicate that there is minimal contamination remaining at the Site. information that is explained in Therefore, EPA is proposing that no further action is

Risk Assessment Report dated

or the environment. This Proposed Plan identifies the all other records utilized

reasoning for no further action and explains the rationale specified in this document are for this preference.

The EPA's decision for no further action represents a Comment Period:

preliminary decision, subject to public review and comment

under Section 117(a) of the Comprehensive Environmental August 24, 1995

Response, Compesation, and Liability Act (CERCLA, known as Superfund), as amended by the Superfund Meeting

Amendment and Reauthorization Act (SARA) of 1986.

August 10, 1995

This Proposed Plan is being distributed to the public in 7:00 P.M.

order to solicit public input. Lewisville High School

Investigation Report dated

by publishing Proposed Plans at Superfund sites, and by public to comment on the Changes to the preferred preferred alternative to comments or additional data decision regarding the elected in a Record of Decision (ROD) consideration all comments request, EPA will extend the

This fact sheet summarizes greater detail in the Remedial July 1995. These documents and by EPA to make the proposal

additional days.

Public July 24,

Thursday,

Public

Date: Thursday,

Time:

Place:

EPA is initiating a thirty (30) day public comment period Richburg, SC

from July 24, 1995 to August 24, 1995, to receive comments on this Proposed Plan, the Remedial comments or call:

Investigation (RI) Report and the Baseline Risk Cynthia Peurifoy

Assessment (BRA) Report. However, EPA will Protection Agency

accommodate requests for informal briefings during the Remedial Branch

week of the Proposed Plan meeting. EPA, in consultation Courtland St, NE

with the South Carolina Department of Health and 30365

Environmental Control (SCDHEC), will select a remedy for 435-9233

Provide written

Yvonne Jones or

US Environmental

North Superfund

345

Georgia

1-800-

contained in the administrative record for this Site. EPA In January 1975, Columbia Organic Chemical Company

and SCDHEC encourage the public to review this up the SEPCO Plant in

information, especially during the public comment period, part of this clean up effort,

to better understand the Site, the Superfund process, and the COCC transported and stored the waste of approximately

intent of this Proposed Plan. The administrative record is 2,000 drums at the Carolawn Site. As payment for services

available for public review during normal working hours, the plant in Clover, South

locally at the site information repository, which is the Carolawn property.

Lancaster County Library, the Chester County Library or in the Record Center at EPA, Region IV's office in Atlanta, Recycling and Disposal, Inc.

Georgia (see page 11).

controlled the site.

permit from SCDHEC for THIS PROPOSED PLAN:

drums containing inert

was given approval to

1. Includes a brief history of the Site, the principle 3-acre fenced portion of the

findings of the RI and a summary of the Baseline SCRDI sold the 3-acre fenced

Risk Assessment;

Carolawn Company.

Clover, South Carolina. As

(COCC) was contracted to clean

rendered during the cleanup of

Carolina, COCC received the

After 1975, South Carolina

(SCRDI), a subsidiary of COCC,

During 1978, SCRDI obtained a

a one-time disposal of 300-400

waste. In October 1978, SCRDI

dispose of empty drums on the

property. After the disposal,

area of the site to the

2. Presents EPA's rationale for its preliminary In 1978, the Carolawn Company began the construction of selection of the preferred alternative; and

With conditional approval of

conducted with one incinerator;

3. Explains the opportunities for the public to incineration never developed. At the

comment on the alternative for the Carolawn site by the Carolawn Company,

(OU2) Superfund Site.

contained a concrete loading dock,

tanks and drums, two

trailers, 14 storage tanks, and as ~-. SITE DESCRIPTION AND HISTORY liquid and solid wastes. An

storage tanks were located The Carolawn Site, located on approximately 60-acres of north. In 1979, SCRDI was land, is an abandoned, waste storage and disposal facility notified by SCDHEC that they would have to clean up the located in Fort Lawn, Chester County, South Carolina. The site is situated less than three miles west of Fort Lawn, and approximately one-half mile south of South Carolina SCDHEC and EPA conducted site Highway 9 (see Figure 1.1). Rural and agricultural areas site. These investigations surround much of the site. The Lancaster & Chester environmental and private residential Railroad and County Road 841 border the site to the south results of these and Fishing Creek borders the site to the east. Wooded

presence of trichloroethane (TCE)

areas and cultivated fields lie to the west and north of the and other solvents in nearby residential wells. The results site.

was contaminated with high

compounds. Due to the The Carolawn site was originally owned by the contamination found and the potential Southeastern Pollution Control Company (SEPCO) of public health and/or the Charlotte, North Carolina. Beginning in 1970, SEPCO used environment, EPA initiated cleanup activities at the site on the site as a storage facility for a solvent recovery plant activities continued located in Clover, South Carolina. SEPCO went bankrupt included removal of in 1974, and abandoned the Site leaving approximately liquid waste from the site.

two incinerators on the site. SCDHEC, a test burn was however, full scale time of abandonment of the the 3-acre fenced area a diked area for storage of incinerators, two storage many as 480 drums containing additional 660 drums and 11 outside the fenced area to the Carolawn site.

During the early 1980's, investigations at the Carolawn included collecting well samples for analysis. The investigations showed the also indicated that the Site levels of metals and organic elevated levels of threat for imminent damage to December 1, 1981. The cleanup through February 1982, and

contaminated soils, drums, and

2,500 drums of solvents on site. SEPCO had been storing Subsequently, in December 1982, the Site was proposed for the drummed solvents in anticipation of incinerating the inclusion on file National Priorities List (NPL). The waste. However, neither an incineration permit nor a the NPL in September, storage/disposal permit was issued to SEPCO by the of local residential wells SCDHEC.

levels of TCE, the Chester

Carolawn Site was finalized on 1983. Since continued sampling showed persistently high

Municipal Sewer District's water main from Highway 9 was Although this area was addressed during an EPA removal extended to four of the five residences living near the site. 1990 field investigation by the These four residents were connected to this alternative Division, some uncertainties water supply in 1985. presence or absence of soil

review of all the available Due to the complexity of the Carolawn Site, and in order to Remedial Investigation and simplify the investigation and response activities, EPA needed to be conducted on OU2 divided the Site into two discrete study areas known as risk assessment and a sound Operable Units (Figure 2). Operable Unit One (OU1) consists of source areas located on a 3-acre parcel within the fenced area of the site and the groundwater located as follows:

OU2). Operable Unit Two (OU2) consists of the land samples from 42 locations located immediately around the fenced area and the land background surface soil sample; located north and west of the fenced area (north and west drum areas).

beneath the entire Site (to include the groundwater beneath

soil samples from 10

one background subsurface On August 29, 1985, a group of Potentially Responsible Parties (PRPs) (the Carolawn Generators Steering Committee) entered into a Partial Consent Decree with the water and 4 sediment samples United States Government to conduct a Remedial that included one background Investigation and Feasibility Study (RI/FS) for OU1. The and one background sediment

action and again during the EPA, Environmental Services still existed as to the contamination. Based on EPA's data, it was determined that a Feasibility Study. (RI/FS) in order to develop a baseline remediation plan.

The RI field activities were

Collected surface soil that included one

Collected 10 subsurface locations that included soil sample; Collected 4 surface from offsite locations

surface water sample

purpose of this RI/FS was to fully characterized the nature and extent of the contamination present at the Site and to identify the relevant alternatives for remedial action. Phase reconnaissance of the Carolawn __.,

I and Phase II of the RI/FS, conducted at the Site between surrounding area in order to identify

1985 and 1989, confumed the presence of volatile organic which are potentially affected

compounds (VOCS) in the groundwater exceeding migration from the site;

Maximum Contaminant Levels ("MCLs") set by the
National Primary Drinking Water Regulations in the Safe
screening to identify

Drinking Water Act. On September 27, 1989, EPA issued threatened species within the site

a ROD for OU1 which selected a groundwater interception performed by contacting

and extraction system as the remedy for groundwater and Wildlife Service. The

contamination at the site. It was also determined that due was collected, reviewed and

to the effectiveness of the removal actions, no source of the investigation.

contamination remained within the fenced area of the site.

However, the findings documented in the ROD for OU1
electromagnetic investigation to

indicated that limited soil data was collected from the west wastes or drums at the site.

and north drum areas located outside the fence; therefore, collection of additional samples was necessary to confirm INVESTIGATION

the Presence or absence of residual soil contamination in these areas. See the Section titled Update on OU1 on page and extent of contamination

10 of this fact sheet for the current status of OU1. defined the potential risks to

environment posed by the Site. A

In response to these concerns, EPA conducted a field total of Fifty-two (52) soil, four (4) surface water, and four

investigation at the Site in 1990. The purpose of the field collected (see Figures 3 and 4).

investigation was to provide additional information on the ROD for OU1 selected a

presence of contaminants in the subsurface soil at the extraction system as the

former storage areas situated outside the fenced area. The remedy for groundwater contamination at the site. Since a

sampling results indicated the presence of VOCs in the soil. groundwater remedy has been selected for the Carolawn

sample;

Conducted a site

site and the

the various habitats

by contaminant

Performed an Ecological

endangered and

area. The screening was

the United States Fish

data from this agency

sununarized as part of

Conducted an

locate any buried

RESULTS OF THE REMEDIAL

The RI investigated the nature
on and near the Site, and
human health and file
total of Fifty-two (52) soil,
(4) sediment samples were
As previously discussed, the
groundwater interception and

site, groundwater was not evaluated in the RI or the to identify the various habitats
Baseline Risk Assessment. All samples collected during the by contaminant migration
RI were analyzed for volatile and extractable organic reconnaissance included rough compounds, pesticides, polychlorinated biphenyls (PCBs), habitat zones present and and metals.

species within each habitat zone.

of species composition or Human Health various habitats were made

However, a baseline ecological The laboratory results for all samples collected were to determine if there is any evaluated to identify compounds that exceeded threshold the environment from previous concentrations (standards) established by EPA and SCDHEC; or were statistically significant compared to background concenwations. These compounds were risk assessment, the laboratory identified as chemicals of potential concern (COPCs). These collected were evaluated to identify compounds were further evaluated to determine the human compounds that exceeded threshold concentrations health risks associated with their exposure to people. The and SCDHEC; or were risks for each of these compounds was estimated in the compared to background Baseline Risk Assessment Report. compounds were identified as

potential concern (ECOPCs). These
The following is a summary of the chemicals of potential
evaluated to determine the
concern identified in each media sampled.
with their exposure to

for each of these compounds
Soil. The results of the surficial soil analyses indicated that
Risk Assessment Report.
there are several COPCs present in the soil cover. These
compounds include: arsenic, barium, beryllium, calcium,
the ecological chemicals of
chromium, copper, iron, lead, magnesium, manganese,
in each media sampled. the
sodium and polyehlorinated biphenyls (PCBs). Other
used to select ecological

the surrounding area it order which are potentially affected from the site. The delineation of the various identification of dominant No quantitative measurements physical characteristic of the during this investigation. risk assessment was performed present or potential risk to site activities. Similar to the human health results for all sample (standards) established by EPA statistically significant concentrations. These ecological chemicals of compounds were further environmental risks associated ecological receptors. The risk was estimated in the Baseline The following is a summary of

potential concern identified

screening criteria that are

concentrations of inorganics and organics were detected in chemicals of potentia concern are specific to ecological

the soil. However, the concentrations of these contaminants receptors; therefore, the COPCs may often include different

were below the typical background concentration ranges for human health assessment.

native soils or were below the threshold standards established by EPA.

surficial soil analyses indicated that

present in the soil cover. These Surface Water and Sediment. There were no COPCs barium, copper, lead, identified for surface water. In addition, no volatile and PCBs, tetrachloroethene, and extractable organic compounds, pesticides or PCBs were toluene. Other concentrations of inorganics and organics detected in any of the Samples. Metals were detected in all However, the concentrations of of the surface water samples. However, the concentrations the typical background of these contaminants were below the typical background native soils or were below the concentration ranges.

The sediment analyses revealed that arsenic is the only of barium, all chemicals chemical of potential concern in sediment. In addition, no eliminated as an ECOPC. volatile organic compounds, pesticides or PCBs were eliminated from sediment during detected in any of the samples. no screening value or

available for compound, Environmental Health to cause a threat to the

established by EPA.

normally precipitates out of Ecological Site Reconnaissance. Black & Veatch personnel and therefore is less conducted a site reconnaissance of the Carolawn site and organisms. it is unlikely that

individual chemicals than the

Soil. The results of the there are several ECOPCs compounds include: arsenic, manganese, mercury, zinc, were detected in the soil. these contaminants were below concentration ranges for threshold standards

Sediment. With the exception detected in sediment were Barium was unable to be the screening process, because background concentration was However, barium is not likely aquatic environment because it solution as an insoluble salt bioavailable to aquatic

terrestrial organisms will come in direct contact with the ingestion of subsurface soil sediment at the site. Therefore, it should be noted that it is (skin) contact with subsurface soil unlikely that barium in sediment will pose a significant risk

Incidental Dermal

carcinogens and non-carcinogens pose different known to bioaccumulate; therefore, this limits the potential health risks, the EPA calculates two possibility that terrestrial as well as aquatic organisms will numbers when estimating health risks: come into direct contact with these contaminants through the food chain. For these reasons, exposure of terrestrial Quotient is calculated for non-carcinogens to and aquatic organisms to barium in sediment was not whether health problems, other than cancer, further evaluated in this Baseline Risk Assessment. associated with a Superfund site. It is derived

to terrestrial organisms at the site. In addition, barium is not

the chemical exposure level at the site by Electromagnetic Investigation. The primary purpose of this level determined to be safe. If the Hazard Electromagnetic Investigation was to locate any buried greater than 1 there may be concern for waste or metal objects at the site. No magnetic anomalies health effects. Hazard quotients are calculated were detected during the investigation. Detection of chemical of potential concern found at the site. magnetic anomalies would indicate the presence of buried overall potential for non-carcinogenic drums.

by more than one chemical, all of the

quotients calculated for each chemical are added SUMMARY OF RISK ASSESSMENT sum of the hazard quotient is called a

Like the hazard quotient, if the hazard CERCLA directs EPA to protect human health and the greater than 1.0 then the contaminants pose a environment from current and potential future exposure to health risk.

hazardous substances at the site. A risk assessment was conducted to evaluate the potential current and future risks is expressed as an incremental probability associated with exposure to the site contaminants. individual developing Cancer over a lifetime as a

exposure to tile potential carcinogen. Human Risk

below summarize the health risks estimated All of the chemicals of potential concern and the media future exposure scenarios.

(soil and sediment) in which these chemicals were found were evaluated in a Baseline Risk Assessment (BRA). A Baseline Risk Assessment is performed at all Superfund sites to determine whether the site poses a current or potential risk to human health and the environment, in

Because types of

different

1. A Hazard
assess
might be

the chemical

by dividing

Quotient is

potential

for each

To assess the

effects posed

hazard

together. The

hazard index.

index is

possible

2. Cancer Risk
Of an

Or an

result of

Tables 1 and 2 for current and

Table 1

absence of any clean-up. Both potential carcinogenic and non-carcinogenic risks were estimated, with respect to POTENTIAL CARCINOGENIC RISK RANGE current conditions and possible future conditions.

Potential human exposure routes (for adults and children)
Exposure 9E-09 - IE-07
evaluated in the BRA included the following:
Pathways

Incidental ingestion of surfical soil

Exposure 9E-09 - IE-05

Dermal (skin) contact with surfical soil

Pathways

Incidental ingestion of sediment from Fishing Creek
Dermal (skin) contact with sediment in Fishing Creek

Unacceptable risks are those which have a Future potential exposure routes for adults and children less than IE-06. No action would be necessary associated with site development that were evaluated further protective of human health if the risk included all of the scenarios listed above in addition to the probabilities are between IE.04 and IE-06. following:

Note:

probability

to be

cottontail rabbit are common terrestrial

upland habitats. Because these two

upland habitats as well as the study

exposure pathway exists to these

Table 9

they were used as surrogates to represent

exposed to contaminated surface soils

POTENTIAL NON-CARCINOGENIC

HAZARD INDEX RANGE

(HQ) method was used to define

Current Exposure 2E-02 - 5E-05

the representative terrestrial receptors

Pathways

pathway. This method involves: 1)

and the easter

species inhabiting

species are common in

area, and a complete

receptors via soil,

the terrestrial species

at the site.

The Hazard Quotient

potential risk to the

via the soil exposure

Current

Future

of each receptor species to Future Exposure 6E-01 - 5E-05

contaminated food and/or soil; 2)

Pathways

scientific studies the highest

roduces no observed adverse effects

exposure level which produces

Note: Unacceptable risks are those

effects (LOAEL) in the representative

which have a hazard index above 1.0

Dividing the estimated receptor species

NOAEL and LOAEL. A LOAEL

is indicative that there may be a

effects on the receptor species.

robin as a potential receptor for the soil

LOAEL HQ values ranged from

the NOAEL HQ values ranged from Carcinogenic risk estimates for current and future accordance with EPA's guidance for conditions are either below the lower limit 1E-6 or within Assessments, remedial goals for the EPA's acceptable range (IE-6 to 1E-4). No nonecological receptors should be bounded by the carcinogenic hazard indices exceeded EPA's acceptable lower end and the LOAEL value on level of 1.0. In summary, EPA has determined that risks to the risk range is between 3.8E-01 to human health from contaminants in the soil and sediment exceed EPA's acceptable level of are within EPA's acceptable risk range and that remediation of the soil and sediment would not be required for the protection of human health. A more detailed discussion of cottontail rabbit, a potential receptor for the exposure routes and presentation of the risk estimates pathway, the LOAEL values ranged from can be found in the Baseline Risk Assessment located in the NOAEL values ranged from the Administrative Record. accordance with EPA's guidance for

Assessments, remedial goals for the Environmental Risk ecological receptors should be bounded by the Estimating the exposure ECOPCs by ingestion of Determining from past exposure level which q (NOAEL) and the lowest observed adverse species; and, 3) exposure level by the based HQ greater than 1 potential for adverse

exposure pathway, th(6.8E-06 to 6.1E-01 and 6.8E-05 to 6.1E+00. In Ecological Risk protection of NOAEL value on the the upper end. Thus, 6.8E-05 which does not 1.0.

Using the american

the soil exposure 9.5E-08 to 6.2E-03 and

Using the eastern

9.5E-07 to 6.2E-02. In

Ecological Risk

protection of

NOAEL value on the

lower end and the LOAEL value on
A qualitative risk assessment was conducted to determine
risk range is between 6.2E-03 to
if ECOPCs posed an unacceptable risk to the ecological
exceed EPA's acceptable level of
receptors on and near the site. All ECOPCs and the media
of concern (surface soil) were evaluated in the ecological
section of the Baseline Risk Assessment.
determined that risks to the

for Drum contaminants in the soil are below
At the Carolawn site, the terrestrial habitats present on the range and that remediation of the soil site property include upland habitats. The american robin for the protection of the environment.

the upper end. Thus, at 9.5E-07 which does not 1.0.

In summary, EPA has
ecological receptors
EPA's acceptable risk
would not be required

PROPOSED FINAL ACTION does not appear to be

local residents.

After careful evaluation of all the exposure routes, estimated carcinogenic and non-carcinogenic health risks, construction of a series of extraction and ecological impacts, the EPA has concluded that the

ground water, a treatment

Carolawn OU2 site does not pose an unacceptable risk to system design to remove contaminants which pose a risk to

human health or file environment. Based on the data environment, and discharge of treated

collected in the RI and the health and environmental risks
The extraction wells have

estimated in the Baseline Risk Assessment, EPA construction of the treatment

recommends that no further action is necessary to provide system are expected to be solicited soon.

additional protection to human health or the environment.

The Baseline Risk Assessment shows no unacceptable to include comments on the

current or future risk for human health from exposure to the water treatment system during

soils or the sediment. The Baseline Risk Assessment shows this comment period. The Final Design, as well as our

no unacceptable risk for ecological receptors from exposure contains a more detailed

to the soils.

water design, can be reviewed at

repositories listed on page 11.

Based on the results of the Remedial Investigation and the Baseline Risk Assessment Report, EPA is recommending no

Owned Treatment Works (POTW)
feasible or acceptable to

This design includes

wells to collect contaminated

system design to remove

human health or the

ground water to Fishing Creek.

been constructed. Bids for

system are expected to be

Final Design for the ground
this comment period. The Final
March 1994 Fact Sheet, which
explanation of the ground
the site information

Individuals should feel free

further action at this site (OU2). However, should future monitoring of the site (e.g. Five-Year Review) indicate that the site poses an unacceptable risk to the environment, then EPA may initiate clean-up actions under the authority of relations program under

CERCLA and in accordance with the National Oil and citizens' concerns and needs for Hazardous substances Pollution Contingency Plan. enable residents and officials of a

the decision-making process.

authorizes technical work on a UPDATE ON OU1 contractors prepare a

based upon discussions

In response to concerns generated by citizens during a leaders and private citizens.

public meeting held on January 10, 1995, EPA collected techniques EPA will use to

two (2) sediment and one (1) surface water sample located communicate effectively with the community during the

within the 3-acre fenced area. All samples were analyzed remedial process. These communication efforts often

for volatile and extractable organic compounds, pesticides, small informal meetings or

polychlorinated biphenyls (PCBs) and metals.

releases, correspondence and

available for review at the site

The laboratory results for all samples collected were evaluated to identify compounds that exceeded threshold concentrations (standards) established by EPA and administrative record and an

SCDHEC; or were statistically significant compared to reports and other documents

background concentrations. In summary all compounds citizens. The administrative record is

were within an acceptable range. The results are available information used by EPA to select

for your review at the site information repository. under the CERCLA. A

at the Region IV EPA Office in

Currently, EPA and the PRPs are continuing to work information repository is a file that

toward implementing the groundwater clean-up for OU1.

such as technical reports and

Based on comments expressed by local residents and the site. The information

officials, EPA is proposing that the Groundwater Treatment repository documents can be reviewed at the library listed

System be constructed as outlined in the Final Remedial below. For information

OPPORTUNITIES FOR PUBLIC INVOLVEMENT

EPA has developed a community Superfund to respond to information as well as to community to participate in Before EPA carries out or site, EPA staff and/or EPA Community Relations Plan(CRP) in the community with local This plan identifies the include telephone contacts, formal public meetings, news fact sheets. The CRP is information repository. EPA establishes an information repository where are made available to a file which contains all a response action for the site duplicate file is maintained Atlanta, Georgia. The contains current information

reference documents regarding

regarding the documents Design, dated November 1992. Discharge to a Publically maintained in the administrative record and information

repository, visit the library listed below or contact file EPA ARE AVAILABLE

TECHNICAL ASSISTANCE GRANTS

community relations coordinator for the site.

To assist communities in

interpreting the technical findings

You are encouraged to visit the information repository and at Superfund sites, communities may apply for Technical

Assistance Grants of up to

contact EPA and SCDHEC representatives listed in this \$50,000. Congress and EPA

document for additional information. EPA would also

have established requirements

for the use of this grant.

Citizens who are interested in

accommodate requests for informal meetings during the a TAG may contact Ms.

public comment period, to further explain the findings of Cynthia Peurifoy at 1-800-435-9233.

tile RI/FS and the Proposed Plan. Individuals interested in arranging briefings should contact EPA's Community Relations Coordinator for the Site.

Administrative Record and Information Repository

Lancaster County Library	Chester
County Library	
313 South White Street	100
Center Street	
Lancaster, SC 29720	
Chester, SC 29706	
(803) 285-1502	(803)
377-8145	

HOURS

HOURS

Monday - Thursday Monday

- Thursday

9:00 am - 8:00 pm 9:00 am

- 7:00 pm

Friday Friday

- Saturday

9:00 am - 5:30 pm 9:00 am

- 5:00 pm

Saturday

Sunday

9:00 am - 5:00 pm

Closed

Sunday Closed

FOR FURTHER INFORMATION

Remedial Project Managers

Operable Unit One - Alfred Cherry
Operable Unit Two - Yvonne Jones
U.S. Environmental Protection Agency
345 Courtland Street, NE
Atlanta, Georgia 30365
(404) 347-7791 or (800) 435-9233

Community Relations Coordinator

Cynthia Peurifoy
U.S. Environmental Protection Agency
345 Courtland Street, NE
Atlanta, Georgia 30365
(404) 347-7791 or (800) 435-9233

Regional TAG Coordinator

Rosemary Patton
U.S. Environmental Protection Agency
345 Courtland Street, NE
Atlanta, Georgia 30365
(404) 347-3931 Ext 6107

South Carolina Project Manager

Richard Haynes
District Engineer
South Carolina Department of Health & Environmental Control
2600 Bull Street
Columbia, South Carolina 29201
(803) 896-4070

GLOSSARY

Administrative Record - A file which is maintained and contains all information used by the EPA to make its decision on the selection of a response action under CERCLA. This file is required to be available for public review and a copy is to be established at or near the site, usually at the information repository. A duplicate file is

maintained in a central location such as a regional EPA and/or state office.

Baseline Risk Assessment (BRA) - An assessment which provides an evaluation of the potential risk to human health and the environment in the absence of remedial action.

Carcinogens - Substances that cause or are suspected to cause cancer.

Comprehensive Environmental Response, Compensation and Liability Act (CERCLA) - A federal law passed in 1980 and

modified in 1986 by the Superfund Amendments and Reauthorization Act (SARA). The Acts create a trust fund, known as

Superfund, from taxes on chemical and petroleum companies, to investigate and clean up abandoned or uncontrolled hazardous waste sites.

Information Repository - Materials on Superfund and a specific site located conveniently for local residents.

National Priorities List (NPL) - EPA's list of uncontrolled or abandoned hazardous wastes sites eligible for long-term clean up under the Superfund Remedial Program.

National Oil and Hazardous Substances Contingency Plan (NCP) - The Federal regulation that guides the Superfund program.

Noncarcinogens - Substances that may cause other adverse health effects besides cancer

Parts Per Million (ppm) - Units commonly used to express low concentrations of contaminants. For example, 1 ounce of

Chloroform in 1 million ounces of water is 1 ppm. If one drop of Chloroforms are mixed in a competition sized swimming

pool, the water will contain about 1 ppm Chloroform.

Potentially Responsible Parties (PRP's) - This may be an individual, a company or a group of companies who may

have contributed to the hazardous conditions at a site. These parties may be held liable for costs of the remedial activities by the EPA through CERCLA Laws.

Public Comment Period - Time provided for the public to review and comment on a proposed EPA action or rulemaking after

it is published as a Proposed Plan.

Record of Decision (ROD) - A public document that explains which cleanup alternative will be used at a National Priorities

List site and the reasons for choosing the cleanup alternative over other possibilities.

Remedial Investigation/Feasibility Study (RI/FS) - Two distinct but related studies, normally conducted together, intended

to determine the nature and extent of contamination at a site and to evaluate appropriate, site-

specific remedies.

Reasonable Maximum Exposure (RME) - A term used in the Baseline Risk Assessment. The RME is the highest exposure

to contaminants that is reasonably expected to occur at a site as is based on the professional judgement of the risk-assessor.

Responsiveness Summary - A summary of oral and/or written public comments received by EPA during a comment period

on key EPA documents and EPA's responses to those comments. The responsiveness summary is especially valuable during

the Record of Decision phase at a site on the National Priorities List when it highlights community concerns for EPA decision-

Safe Drinking Water Act (SDWA) - Federal law passed in 1974 to ensure water supply systems serving the public would meet

minimum standards for the protection of public health. The law was designed to achieve reform safety and quality of drinking

water in the United States by identifying contaminants and establishing maximum acceptable levels.

Superfund Amendments and Reauthorization Act (SARA) - Modifications to CERCLA enacted on October 17, 1986.

Volatile Organic Compounds (VOCs) - Organic compounds which easily change from liquid to a gas when exposed to the atmosphere.

CAROLAWN (OU2) SUPERFUND SITE MAILING LIST COUPON

If you have had a change of address and would like to continue to receive site related information or would like for EPA to add your name and address to the mailing list for the Carolawn (OU2) Superfund Site, please complete this self-addressed form. If you have any questions regarding this mailing list, please call Cynthia Peurifoy at 1-800-435-9233.

NAME:

ADDRESS:

TELEPHONE:() -

USE THIS SPACE TO WRITE YOUR COMMENTS

Your input on the Proposed Plan for the Carolawn (0II2) Superfund Site is important in helping EPA

select a final remedy for the Site. You may use the space below to write your comments, then

fold

and mail. A response to your comment will be included in the Responsiveness Summary,

PROPOSED PLAN PUBLIC COMMENT SHEET

Fold on dashed lines, staple, stamp and mail Name
Address
City State Zip

Cynthia Peurifoy, Community Relations Coordinator North Superfund Remedial Branch/Waste Division U. S. EPA, Region 4 345 Courtland Street, NE Atlanta, GA 30365

United States
Environmental Protection
Agency
Region IV

North Superfund Remedial Branch 345 Courtland Street, NE Atlanta, Georgia 30365

Official Business Penalty for Private Use \$300

Cynthia Peurifoy Community Relations Coordinator (Carolawn NPL SITE)

Attachment B

Public Notices of Public Comment Period and Extension of Public Comment Period

Attachment C

Written Public Comments Received During the Public Comment Period

To: Cynthia Peurifoy, Community Relations Coordinator

North Superfund Remedial Branch/Waste Division

From: Susan K. Helms
Date: August 1, 1995
Topic: Toxic Cleanup

I would like to state my ideas concerning the proposed cleanup of the toxic mess in my community.

- 1. The ground water within a mile radius of the site should be treated with a permanent waste treatment plant which should be built on the site. The treated water should be pumped to the nearest natural creek (Fishing Creek). This plant should be built to allow community to use for future growth after the contaminated water has been treated. The estimated time of completion should be within an eight year period.
- 2. The soil and waste including drums have not been completely cleaned up as reported which was proven at the last public hearing with photos and reports of private citizens. The entire site should be examined again especially for underground drums and further cleanup of the area completed.

Thank you for your support and investigation of the matter. I am concerned for my sons and future grandchildren. I know you would be also if you lived in my community.

South Carolina

Department of Health and Environmental Control

Richard E. Jabbour, DDS

2600 Bull Street, Columbia, SC 29201

Chairman Cyndi C. Mosteller

Brian K. Smith

Rodney L. Grandy

Environment Rodney L (lnmdy

Commissioner: Douglas E. Bryant

Board: John H. Burriss, Chairman

William M. Hull, Jr., MD, Vice

Roger Leaks, Jr., Secretary

Promoting Health, Protecting the

September 19, 1995

John H. Hankinson, Jr. Regional Administrator U.S. EPA, Region IV 345 Courtland Street Atlanta, GA 30365

RE: Carolawn OU-II - Record of Decision

Dear Mr. Hankinson:

The Department has reviewed and concurs with the revised Record of Decision (ROD) dated August 14

1995 for the Carolawn Operable Unit II (OU-II) site. In concurring with this ROD, the South Carolina

Department of Health and Environment Control (SCDHEC) does not waive any right or authority it may

have under federal or state law. SCDHEC reserves any right or authority it may have to require corrective action in accordance with the South Carolina Pollution Control Act. These rights include, but

are not limited to, the right to insure that all necessary permits are obtained, all clean-up goals and

criteria are met, and to take separate action in the event clean-up goals and criteria are not met. Nothing

in the concurrence shall preclude SCDHEC from exercising any administrative, legal and equitable remedies available to require additional response actions in the event that: (1)(a) previously unknown or

undetected conditions arise at the site, or (b) SCDHEC receives additional information not previously

available concerning the premise upon which SCDHEC relied in concurring with the selected alternative;

and (2) the implementation of the remedial alternative selected in the ROD is no longer protective of

public health and the environment.

The State concurs with the selected alternative of "No-Action". The State concurrence on this alternative

is based on the Baseline Human Health Risk Assessment, which determined that the Carolawn OU-II site

does not pose any acceptable current or future risks to buman health. This concurrence is also based

on the Department's above mentioned reservation of rights.

Sincerely, R. Lewis Shaw, P.E. Deputy Commissioner Environmental Quality Control

co: Hartsill Truesdale
 Keith Lindler
 Al Williams, Catawba EQC
 Gary Stewart
 Richard Haynes

Attachment D

Official Transcript of the Proposed Plan Public Meeting

UNITED STATES ENVIRONMENTAL AGENCY

REGION IV

PROPOSED PLAN PUBLIC MEETING

FOR THE

CAROLAWN SUPERFUND SITE

TRANSCRIPT OF PROCEED

AUGUST 10, 1995

RICHBURG, SOUTH CAROLINA

REPORTER:

WACHSMUTH, CVR

P.O. Box 2711 CRS

ROCK HILL, S.C. 29730

(803) 328-9640

REPORTER: SUSAN WACHSMUTH, CVR

Dallas Reporting
VERBATIM COURT REPORTING
P.O. Box 2711 CRS
ROCK HILL, S. C. 29730
(803) 328-9640

1	PROCEEDINGS
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3	United States Environmental ProtectLon Agency
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5	Region IV
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7	Proposed Plan Public Meeting
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11	CAROLAWN SUPERFUND SITE
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13	August 10, 1995 - 7:00 P.M
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15	Lewisville Elementary School
16	Richburg, South Carolina
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20	MS. PEURIFOY - Good evening, everybody. We're
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going to go ahead and get started. I am Cynthia Peurifoy, and I'm the Community Relations Coordinator for EPA, Region IV, with the South Carolina Section of the North Superfund Remedial Branch. I'd like to introduce some people that are here with me tonight: Mr. Al Cherry, who is the Project Manager for Operable Unit One of the Carolawn site; Ms. Yvonne Jones, who is a Project Manager for Operable Unit Two of the Carolawn site; Miss Marlene Tucker, who is our attorney for the site; and Mr. Jan Rogers, who is the Chief of our Section. We also have some people here with us from the South Carolina Department of Health and Environmental Control, Mr. Richard Haynes and Mr. Enayet Ullah.

DALLAS REPORTING Certified Court Reporters Rock Hill, South carolina (803) 328-9640 I'd like to go over with you a little bit our purpose for tonight's meeting. We're going to summarize the remedial investigation and site background, and we're going to talk about the study findings. We're not going to spend a lot of time on the site background tonight because we know that you're really here to talk to us. We're going to summarize the baseline risk assessment and we're going to talk about EPA's preferred cleanup alternative, and we're going to give you the rationale for what we're proposing. We're then going to get into the summary of the groundwater design, and then we're going to ask for your comments, questions and concerns.

~

As you will notice, we have a court reporter here tonight; and we need to make sure that she's able to get down everything that is said, so we're going ask you to come to the microphone, identify yourself, and say whatever you have to say. We're going to also ask you not to interrupt people when they're talking, because she will go crazy, she's already told me. So, let's be very respectful of others and give her a chance to do what she's here to do.

I wanted to talk a little bit about community relations issues. EPA gives Technical Assistance Grants to communities where there are superfund sites.

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This is a grant of fifty thousand dollars to community groups to hire technical advisors. You have to do a 20 percent match, which can be done through in-kind services, cash - - - whatever form you like - - - volunteer services. You must prepare a plan for how you want to use the money, and you can also hire a person to handle the grant for you, an administrator. You cannot use the TAG grant to develop new information or to conduct sampling or underwrite legal actions. The group must be non-profit and must be incorporated.

We have two information repositories set up where you can find administrative records; they are the Lancaster County Library and the Chester County Library. You can also find the groundwater design at the Lancaster library. You have an 800 number, you can

16 call us at any time. It's 1-800-435-9233. Any time 17 you want an update on the site, any time you have any 18 questions or concerns, give us a call. 19 I'm now going to turn things over to Ms. Yvonne 20 Jones. 21 MS. JONES - Basically, as Cynthia stated earlier, 22 due to the fact the majority of, I quess the citizens 23 here - - - please correct me if I'm wrong - - - for the for the 24 most part pretty much have an understanding of what

took place in the background, as far as the Carolawn

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2 MR. CHERRY - Yvonne, they say they can't hear you. Lower your mike a little bit. 3 MS. PEURIFOY - While we're doing this, can 4 5 everybody see the screen okay? 6 MS. JONES - We could also dim the lights if that 7 would be more appropriate. Basically, to summarize, I 8 think, a little bit of the history of the Carolawn 9

site, there was a company by the name of SEPCO Company that, basically, had what we would call a storage and disposal facility that operated on the whole entire site, which we are estimating to be approximately five

acres - - - five to seven acres, I guess which

would be named Carolawn.

history - - - site history.

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Around mid-1970, SEPCO Company basically went bankrupt, and another company by the name of the Carolawn Company basically came in and also operated at the site. Unlike the SEPCO Company, they did not operate on the whole entire site - - - at least to our knowledge they did not operate on the whole entire site but basically they pushed out the drums that were on the inside of the fence, known as SEPCO drums; and, of course, they started their operations within the three acre portion of the fence. I don't know if

everyone can see that. Sometime in 1979, the Carolawn

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Company went bankrupt. Not only that, there was some contamination, basically VOC at this particular time is trichloroethane - - - that was found in a citizen's well. That led to several investigations; mainly SCDHEC, or South Carolina Department of Health and Environmental Control, did a groundwater study. In addition to that, that led to a removal conducted by EPA in 1981.

Basically, as you can see, the areas where the removal took place were, for the most part, within the fenced in area. You can see that to the west portion of the site, which is what we would have considered the West Drum Area and, of course, the North Drum Area.

Basically, this is a photograph taken in 1984. As

you can see, there are still a few horizontal tanks, maybe one vertical tank that is still left on the site. I do not have an aerial photograph that basically shows the site as of this date. However, I can tell you that there is at least one horizontal tank in other words, this is a tank that's fairly large and literally horizontal and, of course, we do have some storage, I guess, drums out there; not really used, I guess, for what they were using them for, but we use them for our remedial investigation activities.

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Basically, the focus of this RI was to study the area on the outside of the fence, and I guess I need to talk a little bit about OU1 and OU2. Based on what I've heard, there seems to be a little bit of concern as to what was considered to be OU2 and what's considered to be OU1. Basically, what we considered OU1 to be was the area within the fence. That is the area where we felt - - or that we know that Carolawn operated on. In addition to that, that also included the groundwater, not only below the three acre fenced in area, but also the groundwater beneath the entire site.

As far as OU2, which is our focus of this investigation that we have currently completed basically, we looked at the West Drum area. I don't know, can everyone see that? We also looked at the North Drum area and, basically, the perimeter around the site. Basically, the area of focus was approximately two acres of land surrounding the chain

- 20 link fence. One of the reasons we did this was - -
- 21 doing Operating Unit One, basically, you know, we
- 22 investigated the area within the fence. However, we
- 23 did not look at the areas right around the perimeter of
- 24 the fence. So, it included that. There was some

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25 concern about whether or not the sediment or the

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surface water had been properly addressed, so we also looked at the sediment and surface water.

Basically, we collected actually 52 soil samples. 42 of those samples were surface soil samples. In other words, those samples were collected from zero to six inches. We collected 10 subsurface oil samples, four surface water samples, four sediment samples. In addition to that, we also conducted what we would call a site reconnaissance to basically determine the type of ecological system that we have out there. We also conducted what we call electromagnetic investigation And, in summary, what that is, it's really - - - it's the way or it's a procedure that we use to determine whether or not we have any buried metal objects below the surface.

And, as you can see here - - - I'm little out of focus - - -but, basically, we've divided the site into what we considered or called grids. Basically, in the areas where we thought there was an indication of contamination, we sampled in a 50 by 50 area, basically, taking composite samples. In the areas where based on the aerial photographs did not really look like they had any - - - I guess, stressed vegetation or indication that there was contamination out there - - - we looked at on a 100 by 100 grid

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- 1 sample. And, basically, this is just a map or a
- 2 figure that shows where we collected the sediment or
- 3 surface water samples. Basically, on all samples taken
- 4 during the focus RI, we ran what we call full scan.
- 5 Basically, what that means is we scan for PCBs,

pesticides, metals, and, of course, all organics and extractable inorganic. EPA and of course, I guess you could say EPA to begin with has what we call a set of screening levels that we consider to be protective of human health and the environment. When we get ready to do our Risk Assessment, basically what we do is we say, do we have any contaminants that are above those screening levels? In addition to that, we

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also take what we call a background sample, where we say, do we have any contaminants that are two times, I guess, whatever our background levels are? If we do have contaminants, we basically sum them up on a list of what we call chemicals of potential concern. Now, I just want to say that does not mean that there's a reason to be concerned. It basically means that, hey, you have some contaminants that are at elevated levels. We don't really know how elevated, we don't even know if there's really a risk that's, you know, been

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generated. But we're going to look at them, in the

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process of using the Risk Assessment, to determine if

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we do have an unacceptable risk. As you can see, there were several inorganics, rather metals, on the surface soil, subsurface soil, and, of course, we had arsenic in the sediment.

Basically, in this particular Risk Assessment, we also looked at the impact or whether or not there was an impact on the ecological receptors out at the site. Again, we also did a list of chemicals of potential concern. I would like to say that, as you can see, this list is not exactly the same as what you would see for human health. The reasons are - - or one of the reasons is because, you know, we are humans and, of course, we're likely to be susceptible or either more or less to certain chemicals than, let's say, a rabbit or a robin. So, that's why you'll see different contaminants of potential concern than what you might see for human health.

Basically, for human health, We looked at several different scenarios. We wanted to see, you know, what would the risk be if someone accidentally - - - I shouldn't say accidentally, but actually ingested

22 surficial soil. What would happen if someone actually came in contact with surficial soil that was 23 24 contaminated by the contaminants that I previously

mentioned. And, as you can see, we looked at what

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2 Fishing Creek or came in contact with the sediment in Fishing Creek. That was what we looked at on the 4 potential exposure routes for adults and children. The 5 reason why I say potential at this particular case is, you know, there's no one living on the site right now. 6 7 However, in the future there could be someone living on 8 the site. So, again, we also have to look at what the 9 chances are of someone ingesting the surficial soil, dermal contact with soil sediment in Fishing Creek, and 10 11 so on. 12 I'm not going to really go into depth on that; however, 13

would happen to someone if they ingested sediment in

at the end of our presentations if you would like to

ask questions, please feel free to do so.

standpoint, we basically looked at the ingestion of either the American robin or, as you can see, the eastern cottontail rabbit for surficial soil. Can everyone see that? It's kind of hard to see. Basically, I guess, it; s the same as what was in the fact sheet, in case you might want to turn there. But, in summary, on the current future exposure scenario, in summary we had maybe, I think it was, nine out of a trillion. The range went nine out of a trillion to one out of one hundred thousand, as far as one being that

Basically, as far as from the environmental

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person that could get cancer if they were exposed to contaminants at the site.

And I guess at this particular point I'd like to explain something. EPA has what we call an acceptable risk range. Basically, that is that one person

cancer from the normal area, so to speak could get cancer out of ten thousand. As you can see, we didn't have anything close to that. We had maybe one person, I think - - - worst-case scenario we had one person out of a million getting, you know, possibly getting cancer. Which, of course, we take action if it's one out of ten thousand. As far as looking at what we call non-carcinogenic risks or, rather, risks that are not cancer causing but are risks, basically EPA has a boundary - - - and that boundary being one - - - at which we would look at taking action. In summary, our worst-case scenario was 0.6. So, again, that's also well below what we would consider unacceptable.

in addition to your additional population getting

Basically, we did the same thing for the American robin and, of course, the cottontail rabbits. It's done a little bit differently because, unlike humans - - - we're basically going out to a spot so many times a day, and that's where they're getting

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their ingestion from. But, in summary, basically those values also fell within EPA's acceptable range.

And, based on the results of the Remedial Investigation, and also Baseline Risk Assessment Report, EPA is recommending no further action at this site, for Operable Unit 2, mainly just because we did not have any unacceptable risk as far as human health or environmental health. However, should future monitoring of the site indicate that the site poses an unacceptable risk to the environment, then EPA - - and I should say EPA in conjunction with the State, of South Carolina - - may initiate clean-up actions.

MR. CHERRY - Hello, I'm Al Cherry, and I'm the Remedial Project Manager for Operable Unit 1. Operable Unit 1 consists of a groundwater clean-up, within the fence, of the site itself - - - in the site itself, right inside the fence. So, for the last couple of years we have been working to complete a design with our consultant, which is Conestoga-Rovers. I think they finally put a good design together, and we hope that this will do the required job.

The Carolawn Groundwater Remediation System consists of two treatment processes. One of them, basically, is an Air Stripping/Clarification Process, and the other process is an activated carbon with a bag

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filter process. The stripping process will be utilized as a primary treatment system, and the activated carbon, the bag filter, will be used as the secondary system. The stripping process is basically designed to handle up to 20 million gallons, and it is an Air Stripping Unit which is basically designed to achieve removal of the particulants of concern. The groundwater system will receive the water from five extraction wells, and these five extraction wells are 120 to 150 feet deep and will go down to what we call bedrock.

Now, there are plans for another addition of five wells, which will extend the capture plume to capture all of the contaminants if it's necessary. These wells are basically designed to produce a flow of one to two million gallons, and we have five of the wells. The treatment system, as I said before, is designed to handle a flow of 10 to 20 million gallons. Each extraction well is four inches in diameter. It has a centrifugal, submergible pump, and it also has a check valve. There's also on each well what they call pressure transducers, and these pressure transducers are installed so they can be incorporated into the automatic system itself. What these pressure transducers will do is control the level of water in

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each one of the wells, and it will tell the pumps when to operate and when not to operate, when to turn on and when not to turn on.

So, now we have the wells. And after the wells we have a tank which is called the equalization tank. Basically, the equalization tank is designed to hold a reservoir from the five pumps, from the five extraction wells. This particular tank will hold up to 475 gallons of water that is extracted from these wells, and basically what this does is it gives us a retention

time. In the tank itself we will have some solids that are being settled out while the water is sitting in the tank, and we have a reservoir so once the controls from the plant itself-decide that the air stripper needs a certain amount of water, then these pumps will kick on and pump water over into the air stripper. This thing is basically designed that they would get a retention of about 23 minutes, that the water coming from the wells would sit and will just set in this tank for 20 full minutes. This will allow some settling of solids that will fall out of this water that initially comes from the well.

The next unit, which is the air stripper, it is designed to handle 20 gallons per minute. Now, we have control valves, and they are located downstream from

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the equalization tank. What they will do is, when told by the control system, they will pump, say, 20 gallons per minute of water to the air strippers Then the way that the air stripper is designed, you actually have water coming in the top and you have air bubbles that are being blown up from a blower from the bottom of this tank, and the contaminated material will, being volatiles, will adhere to the water coming down in the air going up, and you'll get volitization, and your volatiles coming up the top of the stripper.

Now, our consultant will apply for an air permit, if possible, but according to designing some of the calculations, they don't think that we're going to have enough contaminants coming off the top, volatiles coming off the top of this air stripper, to require that the stripper be permitted.

Okay, from that you will get removal of VOCs, as I said before, by the bubbles, and the contaminants attaching to the bubbles and going out the top of the unit itself. And, according to some of the design information that was conducted during the Feasiblilty Study, the Remedial Investigation, from manufacturer's rep we found out that the air stripper will remove 90 percent of the TOC and 45 percent of the DCE and 95 percent of the 1,2,DCE. The air stripper

DALLAS REPORTING Certified Court Reporters Rock Hill, South Carolina (803) 328-9640 has a removal efficiency of 99.6 percent for TCE; 97.5 percent for 1,2,DCE; and 99 percent for 1,1,DCE. These rates, again, are based on the required effluent criteria and expected performance of the equipment itself.

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There are three water probes located in the unit, and basically the water probes will turn the pumps on, turn the pumps off, and give signals to the rest of the system as to the status of the different units. So, the air stripper would discharge the treated material to the clarifier. Now, the second system consists of the bag filter and activated carbon. Now, this could be used two different times. It would be used when you're having some type of maintenance being conducted on an air stripper or when you need to achieve additional removal through the use of activated carbon. Now, the streams are set up so that from the equalization tank the flow goes into the air stripper, you have solids coming off the equalization tank, and you also have the volatile gases going off through the stack of the air stripper itself. Okay, that material from the treated water from the air stripper will then go into a clarifier, and the water from the clarifier can go to the discharge channel and the solids can go to the dry bed, to the sludge drying bed. Or, from

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1 that it can go to the bag filter. The bag filter, what 2 it does basically, it filters solids and keeps from 3 overloading the activated carbon. And so you 4 actually - - - the bag filter has a disposable type 5 filter that you can take out if you need to take it 6 out. You can take it out and you clean it when you get 7 pressure across the filter, or you can run that 8 particular treated material into the activated carbon. 9

Now, the activated carbon has high-efficiency removal, just like the air stripper itself. Very, very seldom will you need to run both of these at the same time, unless you have a failure in one of the systems.

Okay, we have an outfall, and the outfall to Fishing Creek is 1,300 feet. There are several manholes in that particular outfall, where we can take samples and we will know what's going on inside. So, back to the plan itself. We have the watering beds. The sludge from the watering system, it can be disposed into a hazardous-type landfill. We have all these

20 different controls on the system itself which will tell

- 21 us - actually it would take a system, send it back
- 22 to an operator, and we would have pre-set phone numbers
- 23 that if you call it would call the phone number. In
- 24 case of a fire alarm, it would call the phone number.
- 25 In case of a failure in the system, it would call. In

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case you have some type of intrusion, someone interfering with the system.

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24 25 Here, again, this system is also developed so we can have additional extraction wells. So, we have a capture plume now, we have a plume now where it looks as though five wells will capture the contamination that is in the groundwater underneath the site. But, there are several things that can happen. If it's not capturing it, we can build additional wells. We can install additional wells to extend the plume. If it's moving too slowly, additional wells may speed up the capture, and we can look at it that way. So, we think that this system will do the required job, and will do the job that we're looking for.

In future design we actually have on this particular drawing and diagram we have plans if required where we could put in another activated carbon unit along with additional wells. All through this system itself we have what we call sample reports. We can check the efficiency of the unit itself, like we have sample reports after the equalization tank where we can go in and take a sample. And we also have sample reports after the air stripper. Now there is - - which I guess I would probably have to look into it. There are some things that this automatic

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system can monitor, like pH and maybe inorganics, and it will basically tell us if the stream coming into this treatment system itself is chamging. But that's basically the advantage of an equalization tank, so if you've got different wells pumping amounts of different contaminants, once it gets into the equalization tank you will have more of an even flow of - - - even fill, even feed going into the different units itself.

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Now, there have been some questions about whether this plant can handle multiuse uses. I don't know, but our consultants seem to feel that if the money was available and people were interested they maybe could do some additional things to have this. But from what I understand, I don't know if this type system would be able to take municipal sewage or not. I don't think it will, but our consultants are saying it could be possible, but I don't know what the expense will be.

Here, again, we're talking about removal of 90 percent. It would more than reach the criteria - - - our criteria for discharge - - - that we should have less than 26 micrograms per milliliter of trichloroethylene, less than 25 micrograms per liter of dichlorethene, and less than 30 micrograms per liter of total solids. With the filters, with the clarifiers that's located behind the equalization tank, and with

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the bag filter and the activated carbon itself, I don't see how this system could not be able to meet this type of discharge requirements from any of the equipment.

So, that's basically about it. I'll try to summarize this. Again, what we have here, like I said before, we've got proposed now, the system will consist of five extraction wells, and these five extraction wells will handle - - - each well will handle from one to two gallons per minute. We have, then, from that it would be going to an equalization tank, And the equalization tank will hold 475 gallons. From the equalization tank we go to the tray stripper, and the stripper will, by streams of water going up and by streams of water coming down and pumped air going up, the gases will attach to the water bubbles and go out through the stack of the stripper. Then, from that stripper we go to a transfer. We get solids removed from the equalization tank, and they will come down and go to a sand drying filter as they develop in the equalization tank. And then from that, you know, air stripper, and from the air stripper we have a clarifier. There will be retention and settling in the clarifier. The solids from the clarifier can go to the drying bed. Then from the clarifier we will run this material through a bag filter. It would filter again,

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and basically that material would also go down to the dewatering bed. And then from that when you wanted to get additional removal you could run this effluent through an activated carbon filter and from there you could go to your discharge channel and go out into the stream and those solids would also you would take the carbon out and send it to a landfill. So, you've got systems that will guarantee a high removal. So, I'm pretty sure this is going to be a good system. I have drawings in the back if anybody is interested in looking at them. They seem to be real complicated; they're real busy, but they have all of the valves and controls and the control units that operate the plant itself.

 MS. PEURIFOY - Thank you, Al. We have a comment period going now. I missed one of my slides, but it ends August 24th. But it can be extended for an additional 30 days should you make a timely request. I want to do one more thing and we're going to start taking your questions and comments. I want to extend a hearty thanks to two wonderful people, the McMinns who have helped me so much in pulling this together and changing schools and everything. I really appreciate it, guys. Thanks a lot. Public comments, questions?

MS. LISENBY - My name is Donna Lisenby. For those

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of you who don't know me, I have spent some time with the EPA people. I've been asking a lot of questions. I've spent probably over, gosh, a hundred hours studying this particular site. You know, four hours today with the EPA officials answered my questions, so I'm not going to need to ask them any questions tonight. I'm just going to read my comments to the EPA for the record, and I'm just going to sit down.

Comments to the EPA: In your fact sheet in the history section EPA stated that there were a record of 2900 drums dumped on the site. You also stated that 1140 drums were removed. This leaves the total of 1760 drums unaccounted for. You stated that only 7.5 out of

- 14 a total of 82 acres was electromagnetically scanned for
- buried drums. EPA's aerial photos are dated in 1979,
- 16 from which EPA and SCDHEC state there appear to be no
- 17 off-site dumping, however dumping occurred on the
- 18 property from 1970 to 1979. There are no aerial photos
- 19 for 1970, 1971, 1972, all the way to 1978. A dump site
- 20 could have been considerably overgrown by the time an
- 21 aerial photo was flown nine years later. A known
- 22 method of disposal by the operators was to puncture
- 23 barrels, release the contents onto the ground, and
- 24 stockpile empty drums. The only way to determine if
- 25 this occurred on the other 60 acres that remains

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- 1 untested is if all groundwater for the entire site has
- been tested. However, only approximately 10 acres have
- 3 been tested - groundwater has been tested. There
- 4 continues to be physical remains on the property:
- 5 parts of half-buried and rusting and corroded drums;
- 6 green, red, brown and gray unidentified solvents.
- 7 PCBs, solvents, and metals - heavy metals
- 8 were all known site contaminants. These are
- 9 heavyweight contaminants. The operators were clearly
- 10 sloppy and indiscriminate in where and how they dumped.
- 11 Based on these facts, my comments to the EPA are as
- 12 follows: I do not feel a sufficient investigation of
- 13 the entire property has occurred. I think there's
- 14 clear evidence, facts, that could indicate the
- 15 possibility of hidden burial or dump sites somewhere on
- 16 the property. I feel very reassured by the redundant
- 17 and overdesign of the groundwater treatment system.
- 18 However, I feel strongly that the possibility exists
- 19 that further soil, subsoil and other, as yet
- 20 undetected, groundwater plumes could be present
- 21 somewhere on the remaining untested 60 acres.
- 22 Therefore, I would like to ask for further testing of
- 23 the entire site. While this could be goLng on, I would
- 24 also certainly like to see the groundwater get dumped
- 25 and treated with the treatment system that you have

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approved. I'd like to see that started as soon as possible, before the contaminant plume moves into Fishing Creek. As a matter of fact, tomorrow would not have been fast enough for me for groundwater treatment. My third and final comment is that I'd like to commend some of the community members here tonight who would like to see a municipal treatment capability added to the system. I think that's very forward thinking and positive. I support this idea, if the PRP's stated criteria can be met. However, I feel the groundwater needs to be treated as soon as possible. Thank you. MS. HELMS - My name is Susan Helms, H-e-l-m-s, and

MS. HELMS - My name is Susan Helms, H-e-l-m-s, and I am from Richburg, but I teach in Great Falls. I want to thank you, the EPA, and the PRP for allowing the community to give input to your proposal. I've written you, and now I would like to state my recommendations publicly concerning the proposed clean-up of the toxic mess in our community. I feel the groundwater has to be treated on the site immediately, as Donna said, and believe the community deserves a permanent treatment facility. This facility should be built to allow the community to use for future growth after the contaminated water has been treated. The estimated time of completion of the decontaminated water should be within an eight-year period. After hearing from

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residents of the community, I think the oil and the waste, including drums, have not been completely cleaned up as reported. This fact was proven at the last public hearing, with photos and reports from private citizens. The entire site should be examined again, especially for underground drums and further cleanup of the area completed. Thank you for your support and investigation of the matter. I am concerned for my sons and my future grandchildren. I know you would be also if you lived in my community.

MR. NICHOLS - Thank you for giving me the opportunity to speak. My name is Barnett Nichols, I'm on the town council for the town of Richburg. I want to commend the EPA, Mr. Cherry, Yvonne, the whole group, for coming back after January. They really did take a licking and they really got their gall to come back, but we thank them. I like Mr. Cherry's presentation. This afternoon Ms. Yvonne asked me would I go down to the site with her, meet her down there.

- 20 And I told her I would, I'd be there at 5:00. I didn't
- have an opportunity to go over any of it I just 21
- 22 stopped at the gate, but the fence had been changed a
- 23 little bit from when I was down there. In 1979, in
- 24 June, we had a tremendous fire down there and I was a
- 25 respondent, the first respondent, and I know where

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these chemicals went. A lot of them went into the air, 2 a lot of them burned, but a lot of them went into the ground. The reason I asked the lady to put that back 3 on the screen was I want to kind of show you where the 5 fire was. When I arrived, there was a bulldozer in this corner of the site pushing drums, knocking holes 7 in the drums, letting the chemicals out. That's what created the fire. And they had a swell going around 8 9 this way for this runoff. And I have the documentation 10 from the newspaper that they stopped me from pumping 11 the water. And I told them I came down to put the fire 12 out, I was a firefighter. I didn't know anything about 13 hazardous materials, but I was a firefighter. I didn't realize all this-was down there at that time, but it 14 15 was all kind of barrels, five or six hundred lying 16 against the fence. Old drums, you couldn't even pick 17 them up. But at that time there was three 18 trailers sitting ther&. I see two now, but it was three. These - - - there wasn't that many tanks back 19 20 there. I think there was about three or four. But the 21 incinerator had not been built. I don't know what that is, I think it's a barrel site over there, I do not 22 23 know. I have no knowledge of that. But I am 24 confident, I believe that I can stand here and tell you 25 that the drums are gone, because they were recycling

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- 1 the drums that day. That's what they were doing. They
- were letting the chemicals out in the runoff. I can 2.
 - attest to that, that's my belief. I don't believe
- there's any barrels here or anywhere else on the site,

5 I just don't believe it. But I liked the presentation, I liked the stripping, I liked the air filtrs. Are those in tandem? Are both of those in line? 7 MR. CHERRY - Yes.

MR. NICHOLS - Both in line. Are they going to be 10 housed, Mr. Cherry? 11

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MR. CHERRY - Yeah, they have a building to put them in.

MR. NICHOLS - No vandalism can bother them? MR. CHERRY - Well, they're going to have a security system. If someone comes in, it will set it

MR. NICHOLS-I'm totally opposed to a sewer plant. I don't like to see chemicals, I abhor chemicals, you cannot get chemicals out of solids once you put them into solids. And sewage is solids. It would be a costly thing to put into operation a sewer plant down there at this time. Let's get the water out first, and then look at a sewer system. I thank you for letting me speak.

MR. BRUCE - My name is Jim Bruce. I'm a resident

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2 moved to Rock Hill, South Carolina in 1979 from 3 Memphis, Tennessee. We've lived in big cities all over the United States in our 31 years of marriage. We got to Rock Hill in 1979 and we said, man, that's it. 5 We're never going to go nowhere else. We're going to 6 7 stay in Rock Hill the rest of our lives. Well, after 8 about 15 years and the doubling of the Rock Hill 9 population, we said, God, we've got to do it again. 10 So, let's - - - for the sake of quality and for the 11 sake of our grandchildren, let's find a farm to move 12 on. So, we looked. And thanks to brother Jim Gaston 13 (phonetic) back there, he found us a little place below 14 Richburg - - - about five miles below Richburg

of Richburg. It's amazing, you know. My wife and I

and September - - - I'm sorry, August, exactly a year 15 ago, my wife and I moved to that farm. Within three 16 17 months, our son Frank, his wife, and our three 18 grandchildren also moved onto the farm. Within two 19 more months, our daughter Debbie and her husband James 20 moved onto the farm. And within the next 90 days, our 21 son Jimmy and his little girls are going to move on.

22 Well, I was pretty shocked when I heard about 23 Carolawn. I mean, initially it really was no big deal, 24 but then I got wind that it was cited as one of 114

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3 happened since the last meetings. I wasn't - - - I Т 4 missed the meetings because I wasn't aware of it until 5 after - - in fact, that's how I found out about 6 Carolawn. 7 I'm satisfied, from all the conversations that 8 I've had with numerous people, that what EPA has done 9 in that five to seven acres I think is commendable. le. 10 I'm okay with the proposed fix. My only suggestion 11 there is let's get it cranked up, let's get it started, 12 let's get it cleaned up right away. But I've got a real problem emotionally with the of the 60-13 14 plus acres. I have talked to an awful t of people in 15 the community. I genuinely believe from the bottom of my heart that the same gusto that you tested those 16 17 seven acres should be applied to the balance of that 18 60; not only surface testing throughout the 60 acres, 19 but subsurface testing. I believe if you do that - - -20 and please, give us the assurance that there is anything, any contaminants found that those problems 21 22 will be addressed at that time - - - while we are 23 addressing the five to seven acres, let's look at the

up. Well, I started getting involved, as I talked

with some folks. And I really appreciate what's

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balance as well. I want my grandkids to grow up and

not be upset with me because their skin is turning

green, okay. Thank you.

MS. MORRIS - I'm not very mechanically-minded, so speaking for the people who live near the Carolawn site, I didn't quite understand, Mr. Cherry, about this air stripper. One of the things you said, that the

gases would go out of the side of the stripper, does that mean that the pollution that's in the ground is going to be coming out in air form?

MR. CHERRY - Not all of it. Most of it is going to be coming out in solids, that's why I tried to emphasize when you have the groundwater coming out going into the equalization tank, you're going to have solids coming down.

MS. MORRIS - But you will have some air pollution?

MR. CHERRY - Right. But according to some tests
that they've done - - - and they'll have to get a
permit - - it won't be enough to create a problem.

It's below the standard that's allowed to come out of
that unit itself. And the way they're doing this - if you can - - - is I hope - - - I hope that I will be
able to do it, is show that all of these units have to
have a certain retention time. See, it's only 20
gallons per minute that that first unit will take, but
they've got 400 gallons of material in that first tank.
And basically what that's in there for is retention

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time so that you get solids falling out. Then it goes from there and it also will go into, you know, the clarifier. So, you don't have all of this material, you have a very small amount of material coming out of the stack itself.

MR. ROGERS - Can I just clarify a little bit of

that? The air stripper - - - the concentrations in the groundwater, as we talked last time, aren't necessarily what we'd say are real high. They exceed what would be acceptable from long term use as a groundwater source, and that's why we're basically instructed by the law and feel like it's inappropriate to just leave it without trying to remediate. But the concentrations - - - I think we talked last time that when we combine all these fiber covered wells will be about 115 parts per billion, which is, in relative terms, very little. It's not good for long term consumption, therefore we're going to clean it up. But as far as stripping that out in the air, you end up with a very, very low concentration coming out of the airstream, and therefore it doesn't - - we don't expect it would trigger any kind of requirement for a permit or even be regulated as a permitted discharge

24 because the concentration's so low that it wouldn't be 25 deemed to be able to cause any kind of adverse health

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2. MS. MORRIS - Let's hope not. MS. PEURIFOY - Would you give us your name, 3 4 please? 5 MS. MORRIS - I'm Margaret Morris. MS. PEURIFOY - Thank you. 7 MR. ROGERS - Are there any other questions, comments, or statements? 9 MR. NICHOLS - I appreciate the chance to get up 10 here one more time. I would like to see EPA appoint a 11 committee from the community to work with them on 12 setting up this air stripper, or whatever they want to 13 do. I'd like the community to be involved, then we wouldn't have to be wondering what EPA was doing. 14 15 MR. CHERRY - I'd like to get Donna in. I sure 16 would like to have her on my side. Where is she? 17 MR. NICHOLS - We need to be involved with the EPA, 18 the community needs to be involved. 19 MR. CHERRY - Yeah, well, she's been on our side. 20 MR. NICHOLS - Come on up here and give me your 21 opinion. 22 MR. ROGERS - That's, you know, what we talked 23 about this afternoon. From what I understand, we got a

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flavor lastúnight - - - from what Cynthia and Yvonne

talked to the community or some of the members of the

community - - - that they would be more interested in a constant interaction with us. One of the things that was proposed last year in the rewrite of Superfund - - - which never took effect because it wasn't authorized, the bill was never passed, but we as an agency are still looking at it as a tool and implementing it where there's interest ard it's appropriate - - - is what they're calling Citizen Action Groups, which are - - Citizen Advisory Groups, where

- 10 the agency tries to let the community around the site 11 set up a group that fairly represents everybody's 12 interest, that continue to conduct an ongoing dialogue 13 with the agency as we conduct and implement things 14 related to the site. It's envisioned that it would 15 take place earlier than where we are with this site. 16 We certainly don't want to slow things down, we think -17 - - when we were here last time, we certainly heard a 18 lot of opposition to the idea of which discharge option 19 we were talking about then. We've responded to that, 20 come back with the original concept of discharge to the 21 creek, and we seem to get a flavor that people don't 22 necessarily oppose that and would like to go forward 23 with implementing the cleanup. So, I think an area 24 that's real ripe for the Citizen Advisor Group would 25 be this continuation of concern over other areas of
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contamination. It's kind of like taking 60 acres and

2 saying, let's go look for a needle in a haystack. 3 That's - - - I may not have any money to do that. But at least - - - one of the things that's been very effective is you know more about the operations of 6 those sites over the years, because you can run down 7 the people who know something about it or were involved 8 with it or know this, that and the other. If we can come up with some reasonably credible feedback to hone 9 10 in on some areas to explore, we can start focusing in 11 on any concerns about additional contamination at the 12 site. And it's very possible that the Citizen Advisory 13 Group would be a good way to go. So, I guess I'd throw 14 that out for consideration. I think Cynthia, since 15 she's our Community Relations Coordinator, I'm going to 16 talk to her about getting back to y'all and trying to 17 start initiating that. But my one warning is that it 18 can't be a focus group with a predetermined interest. 19 It needs to be a fair representation of the community 20 involved with that group. We would like to do it in a 21 way that it's not too burdensome on people's time, but 22 it does, therefore, keep you more involved with interaction with us about the site. It's one of the 23 24 frustrating things we have is we come and do these

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public meetings - - - partially, as has been pointed

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out earlier, because we're mandated by law - - - but we do feel like we want to interact and find out and can 3 find out things from y'all that we'd have a hard time finding and, you know, identifying in Atlanta. But we 5 generally have low turnout. The Citizen Advisory Group, I think, gets at the issue of creating a more 7 consistent dialogue where you can get better interchange along the way about the life of the site. 8 9 We would hope and believe this is sort of toward the 10 end of the site and that we can get the mplementation 11 and groundwork going. We don't feel like there's any 12 remaining problem at the old Operable Unit Two areas 13 outside of the site, realizing those are focused areas 14 that we're looking at. We can continue to explore the 15 remaining 60 acres if we can start to get an idea of 16 what makes sense to go out and explore. Basically, 17 this will go into an operational phase whereby the pump 18 and treat won't clean up the aquifer quickly. It will 19 have to operate and be monitored and be evaluated over 20 time as we implement that residue. There's a 21 continuing dialogue as to whether we really have the 22 technology to completely clean up the aquifer, but 23 we'll at least implement the existing technology to 24 date to improve it to the point where we feel like 25 we've done everything we can. And we will periodically

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1 evaluate it to determine when are we no longer being effective in what we're doing, and that will be a very 3 good area for the Citizen Advisory Group to be involved in, too. Because if we come back seven or eight years 4 5 from now and tell you we're going to shut it off 6 because we can't do any more, but we haven't cleaned up 7 the aquifer, you're going to be upset, I guess. If we 8 can involve you along the way and get a better 9 understanding of what's technically do-able and what's - - - may, in fact, end up being impracticable. 10 11 We don't want to throw in the towel on the front end,

because we think we can significantly improve that

groundwater. But we don't know that we can accomplish - - absolutely don't know we can accomplish the goal we've set, which is get it back downstream. But that's something I want to throw out to consider, because I think we will pursue the Citizen Advisory Group.

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MS. LISENBY - Of the people in this room, could all the people who reside in this area please stand.

MR. ROGERS - Define the area.

MS. LISENBY - Okay. If you live within a 15-mile radius of the school, could you please stand. Okay. Of those people - - - I'm just trying to, because I know everybody isn't comfortable speaking - - of

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those people, how many of you would like to see further testing of the additional acreage? Could you please also stand if you would like to see further testing. Is there anybody who did not stand up again? Okay. That's about 95 percent of the residents in this room would like to see continued testing of the additional acreage. Thank you.

MR. SMITH - Good evening. My name is Reid Smith, I'm a realtor in Rock Hill. I do a lot of business in the Fort Lawn, Richburg area. I have a question. There's a possibility that at a date that the allowable level of contamination will be raised and then SCDHEC will say, well, this site does not warrant cleanup now because we've raised the level of allowable contaminants. Is that possible? I understand from the last meeting it was right there close, and I think you - - -

MR. CHERRY - I think you were talking about the outside of the fence. Isn't that right?

MR. SMITH - No, the water. The groundwater.

MR. ROGERS - I guess my answer is no. The cleanup goals are established based on health-based evaluations. That comes from toxicology and other disciplines that feed information to the agency. We're

not necessarily fully a health-based agency. Agency

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1 for toxic substances, disease registry, and other 2 people get involved in identifying toxic effects of the chemicals in all environments. We utilize the 3 4 information that comes from that; such as, for 5 carcinogens, a thing called a Slope Factor is established which tends to give you an ability in risk 6 7 assessment basis to weigh the nature of the toxicity of 8 that chemical so that you can incorporate it into the 9 Risk Evaluation at your site. If for some reason additional health studies indicated the Slope Factor 10 was wrong, it could result in a different cleanup goal, 11 12 both lower and higher. That's about the only way you 13 would see a change. The MCLs are basically what we use 14 for groundwater, you know, Maximum Contaminant Levels. 15 Some of those are health-based, some of those are 16 technology-based. When we run risk assessment on some 17 of those, they don't come out to be totally protective 18 in our program, but they're an accepted standard 19 throughout the agency for consumption of that material. 20 It's conceivable those could actually go down. If 21 things change in the future, it changes those numbers. And in rare cases it's conceivable, as we develop 22 23 better knowledge of toxicology, some numbers could go 24 up. But I don't think that's going to be the general

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1 MR. SMITH - On this site. 2 MR. ROGERS - Well, any site. I mean, this is kind 3 of a generic issue. We're using the same toxicology information on this site as we would on any site. 5 MR. SMITH -I realize that. 6 MR. ROGERS - The only difference is the exposure 7 that occurs beneath the site. MR. SMITH - So, it's definitely been cleaned up? MR. ROGERS - Well, yeah, as far as we're 9 10 around - - - as long as we're around and as long as 11 there's a program to be implemented. And the DHEC 12 people we work very closely with who have a corollary program regardless of whether we continue to be around 13 14 or not, if there's anything to pursue, the same issues 15 with the same types of approaches to cleanup. MR. SMITH - Is there a time frame? 16

MR. CHERRY - Can you talk about the bids that went

MR. ROGERS - For this site?

out?

MR. ROGERS - For the cleanup or - -
MR. SMITH - Filter and groundwater. Building the
air stripper and - -
MR. ROGERS - We didn't want to talk about this
because we did want to get feedback on this but the
PRP's basically are ready to out and put this bid on

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the street; and they have, in fact, done it within the last day or two.

MR. CHERRY - Went out on the seventh.

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4 5 MR. ROGERS - And they're, I mean, as anxious to get this thing built and implemented as anybody, since they are doing this, not Federal funds but - - - $\,$

 $\ensuremath{\mathsf{MR}}\xspace$. SMITH - These are the companies that paid before?

MR. ROGERS - Yeah, these are the responsible parties we identified to come in and do the evaluations, the testings, and implement the remedy. They're ready to go and moving forward with the bid process now.

MR. SMITH - And they've explored every possibility? I know going over to Lando was out. What about Fort Lawn? Going down with a pumping station to Fishing Creek and then going back up to - - - if you have to expand on Fort Lawn's treatment system. It would be that much better for the community down there.

MR. ROGERS - They explored some other options, the biggest one being Great Falls. And the sewer line isn't there, and they don't really feel an obligation to lay the sewer lines 10 or 11 miles up to the site. So, I mean, other options as viable as could be perceived were explored.

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MR SMITH - Explored with the idea of maybe some help from local or Federal government to - - - if you had to expand on the sewer treatment sys;em at any time would it be - - - would they explore tha possibility?

MR. ROGERS - They're not opposed to that. The problem is we don't really have any direct authority to

is: what we're building out there for tile material we're going to treat is completely different process-wise than what you would deal with domestic sewage. You're basically pumping water out of the ground that's relatively clear, and the suspended solids basically is a little sand that comes out from the well. Those fall out in the equalization tank, and you basically have clear water at this point that has solubilized contamination. And therefore, the proceeds from there goes fairly simply. Domestic waste systems have to deal with a very large load of solids coming in of a very different nature, and a different treatment process. So, to expand the system basically means to build almost a parallel, totally different process system to deal with domestic sewage.

initiate that kind of a process, and the reality of it

MR. SMITH - They don't have to fill it in on this site; like I say, pump it to somewhere. I know citizens down the road between here and Lando are

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opposed to it.

 MR. ROGERS - Yeah. Generally, you know, there was so much outcry that - - - we don't want it pumped through our neighborhoods to get to the treatment plant, we'd rather see you dump it in the creek. And we have the technology that's available to do that. To reach the standards and monitor it so we feel like it's controllable and safe to discharge to Fishing Creek without causing an undue threat of an exceedance or a significant long term release. So, we have basically what seems to be the most logical place to go with it. And it's the quickest to implement.

MR. SMITH - Right now?

MR. ROGERS - At this point.

MS. TUCKER - I want to clarify something that seems to have caused some great alarm. I'm Marlene Tucker, and I'm the assigned attorney for EPA at this site. And, having had the arduous task of trying to piece together all the facts of how the various owners - - - former owners of the site operated so I could make a case to find who the Potentially Responsible Parties are, I can tell you that the manner in which the former owner, SEPCO, operated at the site was very - - was almost a shuffle game with the waste. In fact, they owned more than one site, so a

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lot of waste went not only to the Carolawn site, but all the sites that EPA has had to cleanup. I just wanted to put some perspective on the disparity between the alleged amount of waste brought to the site - - drums and waste brought to the site - - - versus the record of waste that was taken offsite. A lot of this information is not hard and fast, and it's really hard to put a premium on the estimated amount of waste that the company brought to the site because the records were so sketchy and, in terms of keeping inventory in 11 the 1970's, that wasn't a priority for the company. 12 So, what I really want to stress is that EPA did two 13 thorough removals between '81 and '82 covering the 14 entire site, and I'm pretty confident that all the 15 drums were removed. We have no reason to think there are any drums that weren't disposed of, taken offsite. 16 17 And as Jan had said before, if anyone who lived in the 18 community years and years ago who has any additional 19 information about possible drums on the site, you know, 20 please come forward with that information so we can 21 pursue it. But, as far as EPA is concerned, the site 22 is totally clean as to having any drums, and the removal that was conducted in the '80s took care of any 23 24 drums that were buried or lying around on the site.

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Thanks.

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MR. RAY - I'm Marcus Ray. I'm the mayor of Great Falls. I only have one or two questions for Mr. Cherry. What I would like to know is what the total cost of this operation will be - - - projected total cost of the five-year period. It says here three to five years.

MR. CHERRY - Well, you know, since this is not money - - - maybe someone else at the table will know -- - since this is not money that we are spending - - it's not EPA money. You know, I don't know what the total cost is going to be. What we try to do is make sure that they give us a treatment system that would

13 get the result that we need. So, what we would pay for 14 it, as EPA, private company, you know, we would 15 if we hired Conestoga-Rovers and it cost one thing. If 16 they, a private company, hire them, it doesn't cost 17 half as much as the government. So, I don't know. 18 I've got some estimations that I could reach back 19 and feasibility studies and dig that out. 20 MR. RAY - Where would I find that?

MR. CHERRY - It's in the feasibility study in the records. I can dig that out, but it's just still a rough estimation. And this is an estimation that may be in the millions of dollars, but I have to go back and look. Do you happen to know what that is? I don't

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remember what that is, but - - -

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MR. ROGERS - Let me point out one other sensitivity at this point: PRPs don't particularly like to talk about what they think it will cost while they're in a bidding process. So, you kmow, we have some general numbers from the feasibility study, we have some experience from other operations, we can quess. We tend to caveat our dollars in terms of the whole process of all five years of operation and maintenance and a number of other considerations. It makes it real difficult for us to really pinpoint a number. And really, you know, there's a good reason why the PRPs don't want to flaunt what they think it's going to cost. You know they have - - - you know, for them to bid it, they've got a contractor's estimate right now of what it's going to cost them. But they're not going to disclose that at this point in time.

MR. RAY - In the time span to pump these wells, as I understood it when we were discussing it before, was 30 years or more. Now they're saying three to five.

MR. ROGERS - Yeah. For comparison reasons we sometimes use that 30-year figure and just put all the different considerations and various remedies in an equal light. But, you know, nobody really knows how long this pump and treat will last.

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MR. RAY - This was in our waste water boundary, that's the reason I'm concerned. Another question is operators - - onsite operators. Mr. Cherry said operators. Going to be one operator? Going to be an around-the-clock operator? Is it a fully automated plant?

MR. CHERRY - All that I know is that it would be a certified wastewater treatment plant operator or water treatment operator, and they will have - - it's an automatic system that will run 24 hours. What they will do, they will work it out as to how much time, they want this operator to spend at this site. Now, this will be spelled out before we get into it, and if there's some people that are involved, we would give that information to them.

 $\mbox{MR. RAY}$ - Well, as you and I know, automated equipment can fail.

MR. CHERRY - No, it's not - - - automatic you know, it's not what it says. But, you know, they are supposed to run it and they will have a start up, they will hire people to be there, and it will be determined how much time that this operation is going to have to be there. Because we're also concerned about, like you asked, there's going to be a security problem, too. You put all this equipment out there,

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you just can't leave it out there. People shoot tanks I mean, it happensú

MR. RAY - Wastewater, that's my concern. How much waste we're going to have spilled on the ground before that alarm goes - - - $\!\!\!\!$

MR. CHERRY - Yeah, well, they're going to have to sit down and - - - $\!\!\!\!$

MR. RAY - You people are going to police them closely, is that what you're telling me?

 $\mbox{MR. CHERRY}$ - Well, it will be State, it will be community, it will be all of us.

MR. ROGERS - One of the things that - - - we're having to deal with this in a lot of pump and treat systems related to Superfund on the groundwater, but we're also having to deal with it from the underground storage tank program and other methods or programs

- 17 where they're implementing small, confined pump and
- 18 treats. The technology and the computerization and the
- 19 electronics have evolved to such an extent that

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- 20 basically the people who monitor and deal with operator
- 21 control of wastewater treatment plants are recognizing
- 22 some flexibility as to bring the plant up show me that
- 23 you've got the duplicity and triplicate backups and
- 24 various things that electronically will shut the system

the early phase of the operation of the unit, and we

25 down should something go wrong, and demonstrate it on

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will show some leniency on whether you have to have a 2 full time operator sitting there 24 hours a day to 4 watch a relatively simple operation. So, you know, 5 it's an evolving science at this point. The technology certainly is there that allows it to be done. Yes, 6 there are upsets occasionally where the technology 7 8 fails but, in a relative sense, this water is not toxic 9 directly. The reason we're dealing with it is longterm consumption of this groundwater would be 10 11 considered to be adverse to people's health, so we feel 12 like we should remediate the problem. If I drank a little of it, it's not expected that it would have a 13 14 significantly adverse effect, so a spill would not 15 necessarily immediately cause an adverse effect. We 16 are really addressing the groundwater because if 17 somebody started to use that as a water supply and tried to consume that water for a long period of time, 18 19 we do feel like it would have an adverse effect on 20 their health. 21 MR. HAYNES - It'll have a - - - For example, the

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MR. HAYNES - - - the operator comes there daily

1 flows are and all that. Licensed operator will have

MR. RAY - Do you have a pump - - -

to inspect it and monitor that tells them what the

2 to come in every day and inspect it, make sure

old requirement to monitor - - -

- everything is working. If it shuts down any time an
- 4 operator has to come out before they can even start

5 back up.

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MR. RAY - Wait a minute. I've got the mike. What I'd like to know if you have another plant similar that you could give me a cost on that plant. I'm sure nationwide there must be - - -

MR. ROGERS - Could we do that privately?

MR. RAY - Well, you can write me a letter. She has my card. MS. LISENBY - It's a public meeting and the public's right to know.

MR. RAY - What I'm concerned about also is how widespread is the aquifer under there where these wells are drilled? Are they all the same depth? Are they step drilled in different zones, or what? How widespread is it underground?

MR. ROGERS - We do have experience with - - - let me see if I can figure out what the question is. I guess some discussions took place yesterday about a similar system we have down near Columbia which we implemented. It had some problems that caused the cost to go up a little bit based on problems that occurred along the way, but were not related to the technology

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or anything. But the bid was about two million dollars 1 2 for a system to go in that would - - - I don't know, 3 volume-wise it's probably in a similar size for 4 handling water flow, 10 to 20 gallons a minute. And it was in operation for three or four years, was the 5 6 estimated life of that system. And that factors in 7 subcontractor costs and do you have an operator on the 8 site and do a number of other things. The hardware 9 costs really don't necessarily amount up to that much. 10 The design cost and some of the other labor of 11 continuing to visit the site on a periodic basis - - -12 MR. RAY - Through the years. You says it's 13 between three and five years, taking more than five 14 what would it cost? 15

MR. ROGERS - I don't really have a number for you offhand on that. We can dig up some information for you.

MR. RAY - Much safer and more secure than to have

MR. ROGERS - Well, as we learned last time, there's pros and cons to that argument, but, yeah. MR. RAY - That's all I had.

MS. BRYAN - My name is Nita Bryan, I live in

Edgemoor. I want to thank EPA and congratulate you on

your treatment center that's going in. I have a

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with us today, especially John and Cynthia trucking over there to the property. I got concerned. I wasn't even going to speak, and then I heard again that there are no more contaminants at this site, there's no more drums. And I just want to say that I was there today. Unless someone came in and cleaned them up after I left today, they're still there. And there's materials that are not identifiable in large quantity, and things that neither Yvonne nor Cynthia could tell us what they were. I'm not trying to put y'all on the spot, but they did not know. There were drums under the ground, you could see the edges of them. We stepped on them and pushed them, they appeared to have been there for a long time. So, there's still contaminants of some sort. I don't know what they are, but they are there. And although there have been two site cleanups, as I understand from the report, they're still there after all that groundscraping and all that removal. But my question is that I'm hearing that the community is saying, well, we want you to do continued testing, and that you're in agreement that we could do that and yet, after we talked today, my impression from you was that there really isn't any money left to do that testing. I guess I'd ask you to disclose that to the

comment and a question. One, I thank y'all for going

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community as to whether or not there really is any more finances to support any further testing of the other 60 acres.

MR ROGERS - Currently our status of where we are in funding is sort of questionable. With budget cuts and other things going on there was a recision bill

that was passed this year to basically pull back some monies that were already allocated for 1985 - - - or, 9 1995. Which, at this point in the year, may not have 10 actually been spent anyway. But we did, in fact, as a Region, shut down some starts of some sites in other 11 12 states that were ready to be started, because of that 13 --

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effort to pull back the money and reduce spending for the current fiscal year. As far as we know, we have 15 funds next year. We don't know how long we have them, 16 and we have other dilemmas to deal with. We, like 17 every other Federal agency, have to be appropriated 18 money every year. We, unfortunately, also 19 because of Superfund - - - have the dilemma of dealing 20 with a law that - - - the law really doesn't expire, 21 but part of it does, the part that collects the tax 22 that generates the money to fund the program. That 23 being the case, the program could go on if there's 24 money in the trust fund to continue on, but we don't 25 really know what's going to play out as far as

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reauthorization, when will it be reauthorized and what

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kind of restraints will occur before the reauthorization of Superfund. That being the case, it makes us hard to commit to specifically saying we could get our hands on money to do the kinds of investigation we'd have to do. We certainly would try. And it would certainly be hinged on right now we're going to have to start prioritizing everything to the worst-case-first scenario. That being the case, further investigation of this site might not break out as the worst-casefirst scenario if there are limited. We don't have the answer, that's the bottom line. We would try to get the funds, we would try to go forward, we could do some things in-house of a limited nature with our existing resources in-house, but there are a lot of caveats out there that could impact adversely our ability to continue to do that. If we had some solid leads, we can also work through DHEC to try to pursue some things that way, too. It's an unanswerable question but, I mean, there are options that we would continue to do along. Right now we're not looking total doom and gloom. We think we'll have funds, and we think if there's a legitimate need we can go forward and investigate those things. But it's a little more questionable at this point than it

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typically has been in the past.

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MR. CAMP - My name is Don Camp, and I live in Great Falls. From what I read here about the contaminated water, it's no great risk to anyone right now. Okay. If we're going to treat it and we're going to say it's no great risk, then I'm wondering, are we moving it from Fort Lawn/Richburg area and place it in Fishing Creek and hoping the dilution will do a whole lot. And that's my question. If that be the case, I think we could contain it and dilute it in the Catawba River, and the dilution would be much greater because if you're familiar with Fishing Creek, that water gets about this deep in areas, and all water flows to the south from the area we're in. So, I really think we should think again about putting it in Fishing Creek right now. For the preservation of the southern farms.

MR. ROGERS - Certainly, as we've discussed, we don't intend to dilute it in the creek, we intend to treat it down to acceptable discharge level before that option, just like any discharge for any facility would be required to do.

MR. CAMP - Don't you ordinarily have discharges though that have been fined because they're over limits or - - - I mean, don't EPA and DHEC ordinarily have unauthorized discharges?

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MR. ROGERS - Sure.
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           MR. CAMP - Okay. That's - - -
           MR. CHERRY - But this system is almost what they
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     call an advanced waste treatment system. The carbon
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     and the types of technology - - -
           MR. CAMP - Almost? It is, or almost?
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           MR. CHERRY - Oh, it is. It is.
           MR. CAMP - Well, you said it was almost - - -
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           MR. CHERRY - Yeah, but what I'm saying - - - I
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     hate to say it - - - well, it is when you start talking
     about activated carbon and the type of technology - - -
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MR. CAMP - When you talk of activated carbon, have

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     we found any carbon in the water?
           MR. CHERRY - No, I mean activated carbon to remove
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     the impurities.
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           MR. CAMP - Remove them?
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           MR. CHERRY - Yeah.
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           MR. CAMP - Okay, so it would remain in the
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     carbon - - -
           MR. CHERRY - Right.
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           MR. CAMP - - - - to be placed in another area.
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           MR. CHERRY - Well, it would either be generated or
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     - - - well, yeah, it would.
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          MR. CAMP - Okay. And when you speak of solids - -
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     - we're speaking of water, so when we speak of solids,
     what are we speaking of? Are we speaking of the mud,
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     the sediment, what type solids are we speaking of?
             MR. CHERRY - No, we're speaking of the things that
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     are coming out. Actually, there's some of the
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     contaminants in the solids that's - - - you know, so,
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     actually a lot of this stuff is coming out in the
     solids.
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           MR. CAMP - What type solids?
           MR. CHERRY - Well, the volitus.
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           MR. CAMP - What solids?
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           MR. CHERRY- Well, it's basically only probably
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     five percent, 95 percent water.
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           MR. CAMP - What is it?
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           MR. ROGERS - Suspended solids from the well.
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     Particles from the well.
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           MR. CAMP - Suspended solids?
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           MR. ROGERS - Many - - - now, soil particles from
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     the well, because you're pumping the well and picking
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     up at least, because of the disturbance, some suspended
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     solids related to the material in the well.
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           MR. CAMP - Primarily mud?
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           MR. ROGERS - Not necessarily. Properly installed,
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     it wouldn't be mud.
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           MR. CAMP - Silt?
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 $$\operatorname{MR.}$$ ROGERS - Probably silt and sand $$\operatorname{MR.}$$ CAMP - Silt and sand. Okay, that answered my question.

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CONGRESSMAN SPRATT - I've been asked these questions individually. I'm John Spratt and I'm the Congressman who represents this District. We may as well put it on the record to get some answers. First of all, I think I knew this outfit, SEPCO. In fact, I was about to sue them because they owned the plant up near River Hills.

MS. PEURIFOY - You're talking about Hinson? CONGRESSMAN SPRATT - Yeah. Hinson, is that the name of it now?

 $\ensuremath{\mathsf{MS}}.$ PEURIFOY - Correct. Vaughn Hinson owns the company.

CONGRESSMAN SPRATT - Vaughn Hinson, that's exactly right, yeah. We went there one day because every time it rained these chemical fumes rose from the ground and wafted all over the subdivision and people didn't know what the problem was until I went there with an engineer and we found an Austrian chemist by the name of Behr (phonetic). Maybe you found his name in the records. I remember the guy.

MS. TUCKER - In fact, we tried to locate him. CONGRESSMAN SPRATT - You have found him?

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CONGRESSMAN SPRATT -Well, he was pretty forthright, because I think he was about to be fired; not because he was incompetent, but because I think they were running pretty low on gas at that time. But he explained the problem to us, and it was that they were draining these residues from the North Carolina furniture industry into this particular plant and they were separating out the paint remover from the paint sludge in the paint remover and reselling the paint remover to the furniture industry. And then they had the sludge left over, and they were shipping it down here. He said every time they brought a barrel in - - - a 55-gallon drum in - - - and get it off the back of the truck, and there were no regular means of conveyance, they tended to spill it on the site, all over the site. And then anytime the - - - once they got enough of that stuff spilt over the site it made it pretty slick, and they would call Rock Hill Concrete Company and say send them another load of gravel and

MS. TUCKER - No, we've lost trail of him.

21 they'd just gravel over everything. So, the chemicals

were seeping down into the ground. That was a problem 22

23 up there, but it suggests the kind of way they did

24 business and raises some questions about this site down

25 here. I also happen to have a next-door neighbor who

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running the site down here for a short while. I'm not sure the fire occurred when he was there. His name was was Gregory, maybe you've seen his name somewhere. He told me once a gruesome story of a dog who'd got caught, stuck up in the goo, the viscous mess on the back of this site. He said he almost killed the dog, almost shot him. He was able to get it free, and I think he brought the dog home with him for a time. But that suggested to me that there was a lot of stuff left. The image, the mental picture I have from the way he described it, was that there was this viscous sludge on a good part of the site, enough so that a curious dog wandered into it and got mucked up in it; and he couldn't get out, it was so deep and so sticky. For whatever that's worth, I put that on the record.

was kind of hard up for a job, and he had the job of

Now, I'm interested in exactly, legally, where we stand here. If this is the agreed upon remediation solution, are the PRPs released by court order once this solution is agreed upon as the remediation solution for this site?

MS. TUCKER - Well, there are PRPs known as the Carolawn Steering Committee, comprised of a group of generations that were customers of the Carolawn Company during the time that they owned the site. And their

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1 operating constraints consent decree which has been 2

entered in the District Courts that they have agreed to

conduct the remedy and pretty much build the system.

So, they're operating pursuant to an agreement with EPA.

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CONGRESSMAN SPRATT - Do they then go back to the court and submit to the court this proposed solution, and if EPA finds it to be an adequate solution you sign off on the consent decree and then this becomes the law of the case, there's no further remedy available? If there's water contamination later found on the site, you can't go back to the PRPs?

MR. ROGERS - Let me answer that. This is a unique case. Most Superfund sites, yes, that's the case. I was just alluding that this is a unique case in the sense that we do have a consent decree for this cleanup with the Carolawn group, but it's a partial consent decree. Most sites would have a complete consent decree that takes the site totally to conclusion and provides for covenants and other waivers at the end that you've done everything necessary to deal with this site. The consent decree in place here, because of the nature of the SEPCO operation versus the Carolawn operation - - and they were very much divisible, especially after the drums and waste were pushed

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outside of the fenced area in Carolawn and continued to work just inside of the fence - - - the Carolawn PRPs

made an argument that it wasn't fair for them to do all 3 this work. They would commit to designing and building 4 5 the pump and treat system, so we have a consent decree 6 lodged in court, entered in court, that carries them 7 through to operational activities at the pump and 8 treat. We don't have a consent decree in final form 9 that would settle the long-term operation and 10 maintenance at this site, because they wanted us to 11 bring in the SEPCO parties. And we did bring in the 12 SEPCO parties, and we did have an agreement, and we've 13 actually lodged a consent decree to deal with a joint deal where SEPCO parties - - - some SEPCO parties - - -14 15 and the Carolawn group would continue on with the site, 16 with this activity. Unfortunately, Hinson has now come 17 up, and the SEPCO people over at Hinson are a little 18 upset that they didn't realize they would - - - they 19 think they have double exposure. Nobody knows and can 20 account for where these drums actually ended up but, in fact, their names show up in two places, and they feel 21 22 like that's a little unfair. They've caused us to 23 reevaluate where we are on that lodged consent decree.

24 So, this is still an open issue that we have to deal with through some negotiations and some other matters 25

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being fully funded?

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that relates to Hinson, as well as - - -CONGRESSMAN SPRATT - But if you finally settle 2 3 upon this as the remediation solution for this site, is 4 that final? Does it exonerate them from further 5 responsibility? 6 MR. ROGERS - Only related to the matters at hand. 7 And if this waste is somewhere else on the site, new activity, totally unknown, it's conceivable we have an 8 9 opener in there. But, no, if - - - yes, if they 10 if we found drums right under the site, we'd have a 11 problem. It'd be a fund-lead activity, they'd be 12 exonerated. We don't expect that to be the case, but we've tried to craft that in a narrow enough language 13 14 that it's matters at hand as identified. The public 15 here will want to look at the 60 acres, we've 16 identified the site as originally as a five to seven 17 acre site, so that's really what the investigation has 18 dealt with. I think we would have room to open one of 19 the others, the remaining part of the 60 acres. 20 CONGRESSMAN SPRATT - Now, to what extent does the 21 completion of this task depend upon EPA funding? How 22 much of this comes out of the PRP's pocket for 23 completion? What level of funding - - - to what extent is the consummation of all this dependent upon your 24

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MR. ROGERS - They're paying all the bills, and our 1 2 consent decree has them reimbursing us for our cost to oversee all activities. We have to expend it out-of-3 4 pocket and - - -CONGRESSMAN SPRATT - Yours is mainly an overhead, 5 6 an oversight expense - - -7 MR. ROGERS - Yes. CONGRESSMAN SPRATT - - - and how far you 8 9 10 MR. ROGERS - At this point.

11 CONGRESSMAN SPRATT - Okay. 12 MR. ROGERS - If we trigger other investigations 13 outside of what we know as the site - - -14 CONGRESSMAN SPRATT - You've got to kave the money 15 to undertake that? 16 MR. ROGERS - Yeah. That could be a fund-lead 17 activity, because I think we'll see a little resistance 18 from this group. 19 CONGRESSMAN SPRATT - Yeah. 20 MR. ROGERS - And therefore it does potentially 21 impair our ability to follow through on relevant leads. 22 CONGRESSMAN SPRATT -Well, you were polite enough not to be specific, but the appropriation bill that 23 24 passed the House of Representatives, I did not vote for 25 it, if I can make that clear for the record.

It would

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cut EPA's budget by 37 percent next year, and it would

2 almost certainly have an impact on the conduct of activities like that. Now, let's hope that doesn't 4 pass, but it certainly passed the House of 5 Representatives, so it doesn't indicate that you'll get funded at the President's level of request for the next 6 7 fiscal year. That could truncate some of these 8 activities, that's what I understand you to say. 9 MR. ROGERS - Yeah, our guess is that that's a signal that we think the Senate will moderate a little 10 11 bit, but we guess we're going to get a significant cut. 12 And that cut will impair our ability to deal with 13 everything on the plate, let alone new work. And we 14 don't know what extent that is until we find out what 15 the budget is. We hate a double jeopardy. We have, 16 really, a problem with the reauthorization of 17 Superfund. The appropriation bill specifically - - -18 Superfund says, you can have this money next year, but 19 you can't spend it past December 31st. If that goes 20 through and - - - you know, Superfund's probably a good 21 program to have a confrontation over; let's shut it 22 down for a while and see what happens. In retrospect, 23 in 1986 - - - I've been in this program for 21 years. 24 Emergency response and then this part, more 25 recently - - - in 1986 we suffered some severe damage,

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and probably a two to three year period, because of people who left over an unfunded issue with reauthorization of Superfund. That's concern.

CONGRESSMAN SPRATT - Thank you very for coming.

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MS. JONES - Basically, I'd like to thank everyone for being patient with me, so to speak. And too, Anita, do not worry about putting me down. You are here to address and also - - - not to address but, for the most part, give us your concern. I go out to the site with Anita and Donna today, actually Cynthia and myself, and she is right, we did see I would call tops of drums, so to speak, what I call debris. Something that I did tell her that a lot of times - - - I don't know exactly what it is, but a lot of times when people see - - - this is really interesting to me - - - but a lot of times when people see either drums, or they know it's a Superfund site or, you know, even like I said we have storage tanks out there that we're using for our activities, their perception is that it is hazardous. As I stated before, we have tested the soil. We didn't test the drum tops, you're right. That's something that we're probably not going to do. But I also told her that we do keep open - - - as if, for some odd reason, we do

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know if you want to call it an inquiring, or someone
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     that actually says, I know it's here or we think it's
     here. That's something that we will follow up on. I
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     know I also went out to the site with Mr. Nichols today
     and, you know, he was considered an eyewitness. It
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     sort of, I guess, puts you on the spot; but, again, I
     did not hear that there were buried drums. And really,
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     at all times we would like to keep the channels open.
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     If you know someone who may not be here tonight but
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     who, you know, maybe was there when everything took
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     place, that, you know, has a pretty good feeling that -
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     - - I shouldn't say a good feeling - - - he knows - - -
he knows
     he or she knows that they buried drums - - -
not just
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give what we would call a relevant - - - and I don't

15 anywhere on the site, but they know that those drums were buried or they saw where those drums were buried, 16 17 we would like to talk to them. But, basically, we 18 haven't heard that from the public. We have, you know, 19 people saying that, there's rumors, but we haven't 20 actually - - - we haven't actually had a person that 21 has said, there are buried drums on this site, and this 22 is where they are. We just haven't had that. Again, 23 anybody that you know or that may know something about 24 that, we would be interested in knowing that. But, 25 without that, I mean, we don't really have anything to

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1 follow up on. Thank you.

MS. PEURIFOY - Anybody else have anything to say? We'd like to thank you for coming out tonight. All the comments and questions and concerns that have been raised tonight will be put into a document - - - and it's called the Responsiveness Summary - - - that will part of the Record of Decision. That will be placed in the Information Repository. I will be sending you out a notice when the final decision is made and let you know what's going to happen next. Thank you for coming.

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CERTIFICATE OF REPORTE

State of South Carolina)
County of York)

I, Susan Wachsmuth, CVR, do hereby certify that the aforesaid deponent was placed under oath; that I reported by Stenomask the foregoing proceedings at the time and place herein designated; that my tape was thereafter reduced to typewriting under my supervision; and that the foregoing pages numbered 3 through 68, inclusive, are a true, accurate and correct

transcript of the aforesaid proceedings.

I further certify that I am not a relative, employee, attorney or counsel of any of the parties, nor relative or employee of such attorney or counsel, nor in anyway interested in the event of said cause.

This the 5th day of Sept., 1995, in the City of Rock Hill, County of York, State of South - Carolina.

Susan Wachsmuth, CVR
Court Reporter